

28174

**SEARCH REQUEST FORM**  
Scientific and Technical Information Center

Requester's Full Name \_\_\_\_\_ Examiner #: \_\_\_\_\_ Date: \_\_\_\_\_  
Art Unit: \_\_\_\_\_ Phone Number 30 \_\_\_\_\_ Serial Number: \_\_\_\_\_  
Mail Box and Bldg/Rm Location: \_\_\_\_\_ Results Format Preferred (circle): PAPER DISK E-MAIL

If more than one search is submitted, please prioritize searches in order of need.

Please provide a detailed statement of the search topic, and describe as specifically as possible the subject matter to be searched. Include the elected species or structures, keywords, synonyms, acronyms, and registry numbers, and combine with the concept or utility of the invention. Define any terms that may have a special meaning. Give examples or relevant citations, authors, etc. if known. Please attach a copy of the cover sheet, pertinent claims, and abstract.

Title of Invention: \_\_\_\_\_

Inventors (please provide full names): \_\_\_\_\_

Earliest Priority Filing Date: \_\_\_\_\_

\*For Sequence Searches Only\* Please include all pertinent information (parent, child, divisional, or issued patent numbers) along with the appropriate serial number.

STAFF USE ONLY		Type of Search	Vendors and cost where applicable
Searcher	<i>W.H. Bryant</i>	NA Sequence (#)	STN _____
Searcher Phone #	<i>508-4494</i>	AA Sequence (#)	Dialog _____
Searcher Location		Structure (#)	Questel/Orbit _____
Date Searcher Picked Up		Bibliographic	Dr. Link _____
Date Completed	<i>5/5/08</i>	Litigation	Lexis/Nexis _____
Searcher Prep & Review Time		FulText	Sequence Systems _____
Atticical Prep Time		Patent Family	www/internet _____
Other		Other	Other (specify) _____

KW Cell adhesion; Glycoprotein; Transmembrane; Calcium-binding; Repeat.  
SQ SEQUENCE 1069 AA; 116314 MW; OF3F60C6 CRC32;

Query Match 10.8%; Score 96; DB 11; Length 1069;  
Best Local Similarity 50.0%; Pred. No. 2.17e-02;  
Matches 14; Conservative 8; Mismatches 4; Indels 2; Gaps 2;

Db 887 GIMTVLILLYMMARYCRSKNKGYEA 914  
Qy 31 GILTIVLGVLILLIGCWYCR-R-RNGYRA 56

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RESULT 14  
ID 004557 PRELIMINARY; PRT; 614 AA.  
AC 004557;  
DT 01-JUL-1997 (TRIMBLrel. 04, Created)  
DT 01-JUL-1997 (TRIMBLrel. 04, Last sequence update)  
DT 01-NOV-1999 (TRIMBLrel. 12, Last annotation update)  
DE T7N9.10.  
RA Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.  
OS Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;  
OC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;  
OC Arabidopsis.  
RN [1]  
RP SEQUENCE FROM N.A.  
RA BUEHLER E., DEWAR K., FENG J., KIM C., LI Y., SHINN P., SUN H.,  
RA CONWAY A., CONWAY A., KURTZ D., OJI O., OSBORNE B., SHEN Y.K.,  
RA TORUMI M., VYSOTSKAIA V., YU G., DAVITS R.W., FEDERSPIEL N.A.,  
RA THEOLOGIS A., ECKER J.R.;  
RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.  
DR EMBL; AC00348; AAB61486.1;  
DR PFAM; PF00554; PTRR2; 1.  
SQ SEQUENCE 614 AA; 694.63 MW; 7D875C1F CRC32;

Query Match 10.7%; Score 95; DB 10; Length 614;  
Best Local Similarity 45.8%; Pred. No. 3.19e-02;  
Matches 11; Conservative 5; Mismatches 8; Indels 0; Gaps 0;

Db 564 LIAVIGVNVLYFWYCAHYCYKA 587  
Qy 33 LTVLGVLLIGCWYCRNGYRA 56

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RESULT 15  
ID 061802 PRELIMINARY; PRT; 1510 AA.  
AC 061802;  
DT 01-AUG-1998 (TRIMBLrel. 07, Created)  
DT 01-AUG-1998 (TRIMBLrel. 07, Last sequence update)  
DT 01-NOV-1998 (TRIMBLrel. 08, Last annotation update)  
DE H11E01.3 PROTEIN.  
GN H11E01.3.

OS Caenorhabditis elegans.  
FC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;  
CC Rhabditina; Rhabditoidae; Rhabditidae; Peloderrinae; Caenorhabditis.  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN-BRISTOL N2;  
RX MEDLINE; 94150748.  
RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M., COOPER J., COULSON A.,  
RA BONFIELD J., BURTON J., CORNELL M., COPSEY T., COOPER J., COULSON A.,  
RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAYELLO A., FULLTON L.,  
RA GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L.,  
RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,  
RA LIGHTNING J., LLOYD C., MCNURRAY A., MORTIMORE B., O'CALLAGHAN M.,  
RA PARSONS J., PERCY C., RIFFEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,  
RA SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,  
RA THIERRY-MIEG J., THOMAS M., VAUDIN M., VAUGHAN K., WATERSTON R.,  
RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOLDMAN P.,  
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.  
RL Nature 368:32-38(1994).

**Db** 61 SLHYGTOCALTRRCPEQGFDRDSKVSLOEKNCCEPVVPNAPAYEKLSAEQSPPYSP 118  
 PT used for developing prods. for diagnosis or treatment of expression  
 PT related disorders, partic. melanoma  
 PT Claim 5; Page 14; 26pp; English.  
**Qy** 61 SLHYGTOCALTRRCPEQGFDRDSKVSLOEKNCCEPVVPNAPAYEKLSAEQSPPYSP 118  
 PT This sequence represents the tumour rejection antigen precursor which is  
 CC processed to a tumour rejection antigen presented by HLA-A2 molecules.  
 CC The tumour rejection antigen is not related to tyrosinase. The cDNA  
 CC encoding this sequence was isolated from the melanoma cell line,  
 CC LB-39-MEL. The tumour rejection antigen may be used for diagnosis or  
 CC in vaccines or for therapy of disorders characterised by the expression  
 CC of the tumour rejection antigen precursor, particularly melanoma.  
 SQ Sequence 118 AA;

**RESULT<sup>2</sup>**  
**ID** R84212; standard; Protein; 118 AA.  
**AC** R84212;  
**DT** 20-APR-1996 (first entry)  
**DE** MART-1 melanoma antigen.  
**KW** metastatic melanoma; tumour-associated antigen; melanoma;  
**KW** diagnosis; prognosis; prophylaxis; therapy; immunogen;  
**KW** Mammalian.

**Location/Qualifiers**  
 27 . 47  
**FT** /note= "hydrophobic region"  
**PT** W0929193-A2.  
**PN** 02-NOV-1995.  
**PD** 21-APR-1995; U05063.  
**PR** 22-APR-1994; US-231565.  
**PR** 05-APR-1995; US-417174.  
**PR** (USSH ) US SEC DEPT HEALTH.  
**PI** Kawakami Y, Rosenberg SA;  
**DR** WPI; 95-382963/49.  
**DR** N-PSDB; T02714.  
 PT DNA encoding melanoma antigens recognised by T-lymphocytes - also  
 PT vectors, host cells and antibodies, used to detect, treat and  
 PT immunise animal against melanoma.  
**PS** Claim 11; Page 117; 184pp; English.  
**CC** The melanoma antigen (MART-1) is produced by recombinant DNA  
 CC methods, i.e. preferably using a baculovirus vector for expression  
 CC in insect cell cultures. MART-1 protein is a source of immunogenic  
 CC peptides (see R84136 for peptide M9-2) which are optionally modified  
 CC (see R84783-R84800) and used in medicaments for the treatment or  
 CC prevention (by immunotherapy) of melanoma. Antibodies against MART-1  
 CC and its immunogenic peptides may be used in the detection and  
 CC isolation of MART-1 from a sample, the detection of which is  
 CC indicative of a disease state (melanoma or metastatic melanoma).  
 SQ Sequence 118 AA;

**Query Match** 100.0%; Score 889; DB 1; Length 118;  
**Best Local Similarity** 100.0%; Pred. No. 1.61e-02;  
**Matches** 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

**Db** 1 MPREDAHFIFYGPKKGHHSYTAAEAGIGLTIVGVLLIGCWCRRNGYRALMDK 60  
**Qy** 1 MPREDAHFIFYGPKKGHHSYTAAEAGIGLTIVGVLLIGCWCRRNGYRALMDK 60  
**Db** 61 SLHYGTOCALTRRCPEQGFDRDSKVSLOEKNCCEPVVPNAPAYEKLSAEQSPPYSP 118  
**Qy** 61 SLHYGTOCALTRRCPEQGFDRDSKVSLOEKNCCEPVVPNAPAYEKLSAEQSPPYSP 118

**RESULT<sup>3</sup>**  
**ID** R63158; standard; Protein; 118 AA.  
**AC** R63158;  
**DT** 26-MAY-1995 (first entry)  
**DE** Tumour rejection antigen precursor.  
**KW** Tumour rejection antigen; precursor; HLA-A2 molecule; tyrosinase;  
**KW** isolation; melanoma; cell line; LB-39-MEL; diagnosis; vaccine;  
**KW** therapy.  
**OS** Homo sapiens.  
**PN** W09421126-A.  
**PD** 29-SEP-1994.  
**PF** 09-MAR-1994; U02487.  
**PR** 18-MAR-1993; US-032978.

**PA** (LUDW-) LUDWIG INST CANCER RES.  
**PI** Boon-Falleur T, Brichard V, De Plae E, Traversari C;  
**PI** Van Pel A, Wolfe T;  
**PI** WPI; 94-316544/39.  
**DR** N-PSDB; Q76370.  
**PT** Nucleic acid coding for a tumour rejection antigen precursor - is

PT used for developing prods. for diagnosis or treatment of expression  
 PT related disorders, partic. melanoma  
 PT Claim 5; Page 14; 26pp; English.  
**CC** This sequence represents the tumour rejection antigen precursor which is  
 CC processed to a tumour rejection antigen presented by HLA-A2 molecules.  
 CC The tumour rejection antigen is not related to tyrosinase. The cDNA  
 CC encoding this sequence was isolated from the melanoma cell line,  
 CC LB-39-MEL. The tumour rejection antigen may be used for diagnosis or  
 CC in vaccines or for therapy of disorders characterised by the expression  
 CC of the tumour rejection antigen precursor, particularly melanoma.  
 SQ Sequence 118 AA;

**Query Match** 100.0%; Score 889; DB 1; Length 118;  
**Best Local Similarity** 100.0%; Pred. No. 1.61e-02;  
**Matches** 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

**Db** 1 MPREDAHFIFYGPKKGHHSYTAAEAGIGLTIVGVLLIGCWCRRNGYRALMDK 60  
**Qy** 1 MPREDAHFIFYGPKKGHHSYTAAEAGIGLTIVGVLLIGCWCRRNGYRALMDK 60  
**Db** 61 SLHVGTGTCALTRRCPOEGFDHRSKVSLOEKNCCEPVVPNAPAYEKLSAEQSPPYSP 118  
**Qy** 61 SLHVGTGTCALTRRCPOEGFDHRSKVSLOEKNCCEPVVPNAPAYEKLSAEQSPPYSP 118

**RESULT<sup>4</sup>**

**ID** W00903; standard; Peptide; 21 AA.  
**AC** W00903;  
**DT** 23-MAY-1997 (first entry)  
**DE** Human melanoma MART-1/Aa tumour associated antigen P27-47.  
**KW** Adeno-associated virus; vector; liposome; transfection;  
**KW** melanoma cell; melanoma; MART-1/Aa; adoptive immunotherapy;  
**KW** tumour associated antigen.  
**OS** Homo sapiens.  
**PN** W09703703-A1.  
**PD** 06-FEB-1997.  
**PF** 19-JUL-1996; U12012.  
**PR** 21-JUL-1995; US-001312.  
**PR** 01-NOV-1995; US-007184.  
**PR** 01-DEC-1995; US-566286.  
**PA** (RHON ) RHONE POULENC RORER PHARM INC.  
**PI** Lebkowski JS, Phillip R;  
**DR** WPI; 97-145208/13.  
**PT** Adeno-associated virus-liposome complexes for transfecting dendritic  
 PT cells - for inducing immune response, useful for treating e.g.  
 PT neoplasia or infections  
**PS** Example 5; Page 58; 114pp; English.  
**CC** Tumour associated antigens (W13660-61, W00878-903) can be loaded  
 CC into dendritic cells and used to induce antitumour immunity.  
**CC** Alternatively, the dendritic cells are transfected with adeno  
 CC associated virus plasmid DNA (which includes DNA encoding the  
 CC tumour associated antigen) complexed with cationic liposomes. The  
 CC generate tumour antigen-specific cytotoxic T lymphocytes for use to  
 CC adoptive immunotherapy in a patient having the corresponding  
 CC tumour. A suitable antigen comprises amino acids 27-47 (W00903)  
**SQ** Sequence 21 AA;

**Query Match** 18.7%; Score 166; DB 1; Length 21;  
**Best Local Similarity** 100.0%; Pred. No. 1.8e-06;  
**Matches** 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

**Db** 1 AGIGITIVLGVLLIGCWCY 21  
**Qy** 27 AGIGITIVLGVLLIGCWCY 47

**RESULT<sup>5</sup>**

**ID** W03682; standard; Protein; 291 AA.  
**AC** W03682;  
**DT** 24-SEP-1998 (first entry)  
**DE** Human secreted protein 2.





DE	CD2 ANTIGEN PRECURSOR (T LYMPHOCYTE-LIKE ANTIGEN CD2).		
GN	CD2.		
OS	Mus musculus (Mouse).		
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;		
OC	Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus .		
[1]			
RN	SEQUENCE FROM N.A.		
RX	MEDLINE; 88140313.		
RA	YAGITA H., OKURARA K., NAKAUCHI H.;		
RT	"Molecular cloning of the murine homologue of CD2. Homology of the molecule to its human counterpart T11.";		
J.	Immunol. 140:1321-1326(1988).		
RL	EMBL; M18934; AAA37397.1; -.		
DR	HSSP; P08921; 1ATB.		
DR	MGD; MGI:38320; Cd2.		
KW	Caenorhabditis elegans.		
FT	SIGNAL 1 17 POTENTIAL.		
FT	CHAIN 18 344 POTENTIAL.		
SEQUENCE	344 AA; 383337 MW; AFEA9175 CRC32;		
Query Match	Score 99; DB 11; Length 344;		
Best Local Similarity 40.4%	Pred. No. 6.72e-03;		
Matches 21; Conservative 9; Mismatches 18; Indels 4; Gaps 4;			
Db	198 PEKGLSF-YVTVGVGAG-GLLVLL-VALEIFC-TCKRKRKRNRRKDEELEI 245		
Qy	13 PKKGHGHSYTAAEAGIGLIVLGSVLLIGCWCRRLNGYRALMDKSLEV 64		
RESULT	3	PRELIMINARY;	PRT; 623 AA.
ID	061391		
AC	061391;		
DT	01-AUG-1998 (TREMBLrel. 07, Created)		
DT	01-MAY-1998 (TREMBLrel. 07, Last sequence update)		
DT	01-MAY-1999 (TREMBLrel. 10, Last annotation update)		
DE	GLY5A (EC 2.4.1.41) (POLYPEPTIDE N-ACETYLGLACTOSAMINYLTRANSFERASE)		
DE	(PROTEIN-UDP ACETYLGLACTOSAMINYLTRANSFERASE).		
GN	GLY 5.		
OS	Caenorhabditis elegans.		
OC	Eukaryota; Metazoa; Nematoda; Rhabditida; Rhabditina; Rhabditidae; Rhabditina; Rhabditida; Caenorhabditis.		
[1]			
RN	SEQUENCE FROM N.A.		
RX	STRAIN=N2;		
RX	MEDLINE; 98192520.		
RA	HAGEN F. K., NEHRKE K.;		
RT	"cDNA Cloning and expression of a family of UDP-N-acetyl-D-galactosamine:polypeptide N-acetylglactosaminyltransferase sequence homologs from Caenorhabditis elegans."		
RT	J. Biol. Chem. 273:8268-8277(1998).		
RL	-1 - CATALYTIC ACTIVITY: UDP-N-ACETYL-D-GALACTOSAMINE + POLYPEPTIDE - UDP + N-ACETYL-D-GALACTOSAMINE.		
CC	-1 - COFACTOR: MANGANESE; ALCTUM.		
DR	EMBL; AF01835; AAC13671.1; -.		
DR	PFAM; PF00535; Glycos_transf_2; 1.		
DR	PFAM; PF00652; Ricin_B_lecitin; 1.		
KW	Transferase; Glycosyltransferase.		
SEQUENCE	623 AA; 71014 MW; C061803B CRC32;		
Query Match	Score 99; DB 5; Length 623;		
Best Local Similarity 24.1%	Pred. No. 6.72e-03;		
Matches 19; Conservative 20; Mismatches 38; Indels 2; Gaps 2;			
Db	9 ILKVLLVPFWICSLIF-PAATSNDSSQIGGNNDLANKIAEANHPKAAKDVQGFGP 67		
Qy	36 ILGVLLIGCG-WYCRRGYRALMDKSLVHQTCALTRRCQEGFDHRDKSVLSQEKNCE 94		
Db	68 PIEPEPVVNNKVVEEQP 86		
Qy	95 PVVNPAPPAYEKLSAEQSP 113		
RESULT	5	PRELIMINARY;	PRT; 626 AA.
ID	061392		
AC	061392;		
DT	01-AUG-1998 (TREMBLrel. 07, Created)		
DT	01-AUG-1998 (TREMBLrel. 07, Last sequence update)		
DT	01-MAY-1999 (TREMBLrel. 10, Last annotation update)		
DE	GLY5B (EC 2.4.1.41) (POLYPEPTIDE N-ACETYLGLACTOSAMINYLTRANSFERASE)		
DE	(PROTEIN-UDP ACETYLGLACTOSAMINYLTRANSFERASE).		
GN	GLY 5.		
OS	Caenorhabditis elegans.		
OC	Eukaryota; Metazoa; Nematoda; Rhabditida; Rhabditina; Rhabditidae; Rhabditina; Rhabditida; Caenorhabditis.		
[1]			
RN	SEQUENCE FROM N.A.		
RX	STRAIN=N2;		
RX	MEDLINE; 98192620.		
RA	HAGEN F. K., NEHRKE K.;		
RT	"cDNA Cloning and expression of a family of UDP-N-acetyl-D-galactosamine:polypeptide N-acetylglactosaminyltransferase sequence homologs from Caenorhabditis elegans."		
RT	J. Biol. Chem. 273:8268-8277(1998).		
RL	-1 - CATALYTIC ACTIVITY: UDP-N-ACETYL-D-GALACTOSAMINE + POLYPEPTIDE - UDP + N-ACETYL-D-GALACTOSAMINE.		
CC	-1 - COFACTOR: MANGANESE; CALCIUM.		
DR	EMBL; AF01836; AAC13672.1; -.		
DR	PFAM; PF00535; Glycos_transf_2; 1.		
KW	Transferase; Glycosyltransferase.		
SEQUENCE	626 AA; 71383 MW; BB318A9D CRC32;		
Query Match	Score 99; DB 5; Length 626;		
RESULT	4		

Best Local Similarity	Matches	Pred.	No.	6.72e-03;				
Local Similarity	19;	Conservative	20;	Mismatches	38;	Indels	2;	Gaps
Db	9	ILKLVLLPVWICSLIF-PAATSDNSQTSIGNSNNFLANKIAEANHPKAQDVIQFGFP	67					
Qy	36	ILGVLLIGG-WYCKRRNGTALRMLDKSLHVGTCQALTRRCQPEGDHRSKVSLOERKNE	94					
Db	68	PIEPEPVENNKVEEEQF	86					
Qy	95	PVVPNAPPAYEKLSAEQSP	113					
RESULT								
ID	060245	PRELIMINARY;		PRT;	1069	AA.		
AC	060245;							
DT	01-AUG-1998	(TREMBBLE).	07,	Created				
DT	01-AUG-1998	(TREMBBLE).	07,	Last sequence update				
DT	01-NOV-1999	(TREMBBLE).	12,	Last annotation update				
DE	PCDH1A	(BH-PCDH1A).						
OS	Homo sapiens (Human).							
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;							
OC	Eutheria; Primates; Catarrhini; Hominidae; Homo.							
[1]								
RN		SEQUENCE FROM N.A.						
RP		YOSHIDA K.; YOSHITOMO-NAKAGAWA K., SEKI N., SASAKI M., SUGANO S.;						
RA		Genomics 0;0-0(1998).						
RL		-!!- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN (BY SIMILARITY).						
CC		EMBL: AB006755; BAA25194.1; - .						
DR		HSSP; P15116; INCJ.						
DR		PROST; BS00232; CADHERIN; 6.						
DR		PF00028; cadherin; 6.						
DR		PRINTS; PRO05; CADHERIN.						
DR		Cell adhesion; Glycoprotein; Transmembrane; Calcium-binding; Repeat.						
KW		SEQUENCE: 1069 AA. 116104 MW. E173200 C032.						

Query Match	11.18;	Score 99;	DB 4;	Length 1069;
Best Local Matches	Similarity 50.08;	Pred. No. 6.72e-03;		
Matches 14;	Conservative 8;	Mismatches 4;	Indels 2;	Gaps
Db	887 GIMTVLILIVVMARYCRSKNKNKNGYE A 914  :     :  :: :    : :			
Qy	31 GILTVLGVLLGGWCRR-RNGYRA 56			
RESULT				
ID	060246	PRELIMINARY;	PRT;	1072 AA.
AC	060246;			
DT	01-AUG-1998 (TREMBLrel. 07, Created)			
DT	01-AUG-1998 (TREMBLrel. 07, Last sequence update)			
DT	01-NOV-1999 (TREMBLrel. 12, Last annotation update)			
DE	PCDH7 (BH/PCDH) B.			
OS	Homo sapiens (Human)			
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;			
OC	Eutheria; Primates; Catarrhini; Hominidae; Homo.			
RN	[1]			
SEQUENCE FROM N.A.				
RA	YOSHIDA K., YOSHINO-MAKAGAWA K., SEKI N., SASAKI M., SUGANO S.;			
RL	Genomics 0:0 (1998).			
CC	-1. SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN (BY SIMILARITY).			
DR	EMBL: AB006756; BDR: BAR25195.1; -.			
DR	HSPP: PI5116; INCJ: -.			
DR	PROSITE: PS00232; CADHERIN: 6.			
DR	PFAM: PF0028; cadherin; 6.			
DR	PRINTS: PR00205; CADHERIN.			
KW	Cell adhesion; Glycoprotein; Transmembrane; Calcium-binding; Repeat.			
SQ	SEQUENCE 1072 AA; 116463 MW; A3DF367C CRC32;			
Query Match	11.18;	Score 99;	DB 4;	Length 1072;
Best Local Matches	Similarity 50.04;	Pred. No. 6.72e-03;		
Matches 14;	Conservative 8;	Mismatches 4;	Indels 2;	Gaps
Db	887 GIMTVLILIVVMARYCRSKNKNKNGYE A 914  :     :  :: :    : :			

31 GILTIVLGLLIGCWCR-R-RNGYRA Qy						
RESULT	ID	PRELIMINARY;	PRN;	PRN;	PRN;	PRN;
8	060247					
AC	060247;					
DT	01-AUG-1998	(TREMBLrel. 07	Created)			
DT	01-AUG-1998	(TREMBLrel. 07	Last seqn			
DT	01-NOV-1999	(TREMBLrel. 12,	Last anno			
DE	PCDH7	(BH-PCDH)C.				
OS	Homo sapiens	(Human).				
OC	YOSHIDA K., YOSHITOMO-NAKAGAWA K., SIRI					
QC	Genomics 0:0-0(1998).					
RN	[1]					
SEQUENCE FROM N.A.						
RA	YOSHIDA K., YOSHITOMO-NAKAGAWA K., SIRI					
RL	Genomics 0:0-0(1998).					
CC	[1]-SUCCESSIONAL LOCATION: TYPE I MEM					
DR	EMBL: AB006757; BAA25196.1; -.					
DR	HSSP: P15116; INCHU.					
DR	PROSITE: PS00232; CADHERIN_5.					
DR	Pfam: PF00028; cdherin_5.					
DR	PRINTS: PR0205; CADHERIN.					
KW	Cell adhesion; Glycoprotein; Transmembrane					
SQ	SEQUENCE: 1200 AA; 130337 MW; 56P.					
Query Match	11.18;	Score 99				
Best Local Matches	Similarity 50.08;	Pred. No				
14;	Conservative	8;	Mis			
Db	840 GIMTVLILLITVMARYCRSKKNKGYEA 867					
Qy	:    :  :   :    :    :    :					
31 GILTIVLGLLIGCWCR-R-RNGYRA 56						

RESULT	9	PRELIMINARY;	PRT;	
ID	057537	RP		
AC	057537;	SEQUENCE FROM N.A.		
DTP	01-JUN-1998	BRADLEY R.S., ESESETH A., KINTNER C.		
DTP	01-JUN-1998	RNA		
DTP	01-NOV-1999	CHORDATA; CRANIA		
DEF		Eukarya; Metazoa; Chordata; Crania		
GNP		Batrachia; Anura; Mesobatrachia; Pipe		
OS	Xenopus laevis (African clawed frog)	Xenopus.		
OC		[1]		
OC		RP		
OC		SEQUENCE FROM N.A.		
DR	RA	BRADLEY R.S., ESESETH A., KINTNER C.		
DR	RA	RNA		
DR	CRC	CHORDATA; CRANIA		
DR	CRC	Eukarya; Metazoa; Chordata; Crania		
DR	EMBL	Batrachia; Anura; Mesobatrachia; Pipe		
DR	EMBL	Xenopus laevis (African clawed frog)		
DR	HSSP			
DR	HSSP			
DR	PROSITE			
DR	PROSITE			
DR	Pfam			
DR	Pfam			
DR	PRINTS			
DR	PRINTS			
KW	KW	Cell adhesion; Glycoprotein; transmembrane protein		
SQ	SQ	SEQUENCE 1035 AA; 113713 MW; 7E41		
		Query Match 11.0%	Score 98	
		Best Local Similarity 50.0%	Pred. No 8;	
		Matches 14; Conservative	Miss 8;	
Db	859	GIMTVLLILVVVMARYCRAKSKNGYE 88		
		:     :   :   :     :       :		
Qy	31	GILTIVGVLILLIGNWYCR-R-RNGYRA 56		
RESULT	10	PRELIMINARY;	~	PRT;*
ID	P93050	RP		
AC	P93050;	SEQUENCE FROM N.A.		
DR	01-MAY-1997	BRADLEY R.S., ESESETH A., KINTNER C.		
DR	01-MAY-1997	RNA		
DR	PROSITE	CHORDATA; CRANIA		
DR	PROSITE	Eukarya; Metazoa; Chordata; Crania		
DR	PRINTS	Batrachia; Anura; Mesobatrachia; Pipe		
DR	PRINTS	Xenopus laevis (African clawed frog)		
DR	HSSP			
DR	HSSP			
DR	PROSITE			
DR	PROSITE			
DR	Pfam			
DR	Pfam			
DR	PRINTS			
DR	PRINTS			
KW	KW	Cell adhesion; Glycoprotein; transmembrane protein		
SQ	SQ	SEQUENCE 1035 AA; 113713 MW; 7E41		
		Query Match 11.0%	Score 98	
		Best Local Similarity 50.0%	Pred. No 8;	
		Matches 14; Conservative	Miss 8;	

	Matches	15;	Conservative	9;	Mismatches	6;	Indels	3;	Gaps	3;
01-MAY-1997 (TREMBLrel. 03; Last sequence update) 01-NOV-1999 (TREMBLrel. 12; Last annotation update)	Db	263	EPCGIALVLLILAVVILYIWYRRKNSYK	295						
RKF3 OR T9U23_16.	Qy	26	EAAGIGLTV-ILGVLLIGC-W-YCRRGYR	55						
Arabidopsis thaliana (Mouse-ear cress). Eukaryota; Viridiplantae; Streptophytina; Embryophytina; Tracheophytina; euphyllophytes; Spermatophytina; Magnoliophytina; eu dicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.	RESULT	12	PRELIMINARY;	PRT;	604 AA.					
[1]	ID	004098;								
SEQUENCE FROM N.A. STRAIN=COLUMBIA C24; KEPFBUNDTE S., LINACERO R., ROUZE P., GALIS I., MACAS J., DEBOECK F., HERNALSTEINS J., DE GREVE H.; Submitted (JAN'1997) to the EMBL/GenBank/DBJ databases.	AC	004098;								
[2]	DT	01-JUL-1997 (TREMBLrel. 04; Created)								
STRAIN=COLUMBIA; TAKAHASHI T., MU J.-H., GASCHE A., CHUA N.-H.; Submitted (OCT'1997) to the EMBL/GenBank/DBJ databases.	DT	01-NOV-1999 (TREMBLrel. 12; Last sequence update)								
[3]	DE	RECEPTOR-KINASE ISOLOG (FRAGMENT).								
SEQUENCE FROM N.A. STRAIN=CV. COLUMBIA;	GN	T20D16_21.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	Arabidopsis thaliana (Mouse-ear cress). Submitted (FEB'1999) to the EMBL/GenBank/DBJ databases.								
EMBL; Z84402; CAB06335_1;	RA	DR	U95913; AAC65490_1; -							
EMBL; AF024650; AAC5045_1;	RA	PFAM; PF0050; LRR; 4;								
EMBL; AC006072; ADI3705_1;	RA	PFAM; PF00059; Kinase; 1.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					
Query Match	10	8%;	Score 96;	DB 10;	Length 604;					
Best Local Similarity	51.7%	;	Pred. No. 2.17e-02;							
Matches	15;	Conservative	4;	Mismatches 9;	Indels 1;	Gaps 1				
[1]	RP	SEQUENCE FROM N.A.								
STRAIN=CV. COLUMBIA;	RC	SEQUENCE FROM N.A.								
LIN X., KAUL S., SHEA T. P., FUJII C.Y., SHEN M., VANAKEN S.E., BRANDON R.C., BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITON M., CARRERA A.J., CREALY T.H., BUELL C.R., TOWN C.D., NIERNAN W.C., FRASER C.M., VENTER J.C.;	RA	SEQUENCE FROM N.A.								
EMBL; Z84402; CAB06335_1;	RA	SEQUENCE FROM N.A.								
EMBL; AF024650; AAC5045_1;	RA	SEQUENCE FROM N.A.								
EMBL; AC006072; ADI3705_1;	RA	SEQUENCE FROM N.A.								
DR	NON_TER	FT	1							
DR	SEQUENCE	SQ	604 AA;	66816 MW;	64A9194F CRC32;					



DE Tumour antigen booster peptide Melan-AMART-1 HLA-A2 #2.

KW - Tumour antigen; booster peptide; immune response modulation; allergy;

KW immune response enhancer; tumour cell; tumour rejection antigen;

KW leukocyte antigen presenting molecule; autoimmune disease;

KW allograft rejection.

DS Homo sapiens.

DN #09858956-a2.

PD 30-DEC-1998.

PD 19-JUN-1998. U12894.

PR 23-JUN-1997. US-080979.

PR (LUDWIG) LUDWIG INST CANCER RES.

PA Boon-Faillon T, Uttenhoffen C, Warnier G;

DR WPI: 99-105612/9.

PT Immunization methods using viruses expressing antigen for priming and booster immunizations - useful for modulating immune responses against antigen, e.g. enhancing immune response against tumour cells expressing tumour rejection antigens

PT disclosure; Page 10; 33PP; English.

RS This sequence represents a tumour antigen booster peptide that can be used in the method of the invention. The method is for modulating immune response in a mammal against an antigen, and comprises:

CC (A) inducing an immune response by: (i) administering a virus containing a nucleic acid molecule encoding the antigen or its precursor to generate an immune response; and (ii) administering at least one booster dose comprising a peptide including the antigen, in an adjuvant, in a combination effective to enhance the initial immune response; or

CC (B) reducing an immune response as defined for (A) but using a non-adjuvant with the peptide which includes the antigen in an amount effective to reduce the initial immune response. Method (A) is used to enhance the immune response against tumour cells expressing tumour

CC rejection antigens, and against pathogens in subjects having human

CC leukocyte antigen presenting molecules. Method (B) is used to reduce the immune response in allergy, autoimmune disease, and allograft rejection.

CC Method (A) provides an immunisation method which, unlike prior art, is not limited by the host immune system.

Query Match Score 56: DB 1: Length 0.

Best local similarity 100.0%,  
FNU. NO. 1.00/0.1;  
Matches 9; Conservative  
0; Mismatches 0; Indels 0;  
Gaps

	Matches	Mismatches	Indels	Gaps	O:
9; Conservative	180	105	10	10	0
9; Strict	199	1205	10	10	0
9; lenient	199	1205	10	10	0
9; lenient, DR	199	1205	10	10	0
9; lenient, DR, PII	199	1205	10	10	0
9; lenient, DR, PII, RTT	199	1205	10	10	0

卷之三

PI or antigen in the lymphatic system sustained CMI response used to +

The present invention describes a disclosure, made on 4th, 1999, in English.

RESULT 3

CC method comprises: (a) delivering RESULT

CC the level of the antigen in the m

e.g. a differentiation antigen, a CC

immunisation; tumour; infectious disease; hepatitis; AIDS malignant melanoma; viral disease; hepatitis; cancer;

gene antigen; or a viral antigen such as cancer gene; melanoma; viral disease; hepatitis; AIDS; immunosuppression; tumour; infectious disease; immunotherapy; cancer; malignant melanoma; viral disease; hepatitis; AIDS; immunosuppression; tumour; infectious disease; hepatitis; immunotherapy; cancer;

Homo sapiens.

e.g. viral disease such as rubella can provide to the lymphatic system

PD 21-JAN-1999.

CC that is necessary to keep CTL active  
CC place all the memory on one side of the bus

PR 10-DEC-1997; US-988320.

CC antigens given in the present inventio

PA (CTL1<sup>-</sup>) CTL IMMUNOTHERAPIES CORP.

PA (CTL-1) CTL IMMUNOTHERAPIES CORP.

WFR 1991-120514/10: Production of cuttings from seabed - by methods

Inducing a cutaneous lymphocytic reaction by means of prednisone.

antigen, the lymphatic system of man so as to provide a sustained CTL response used to treat AIDS.

1 AND CRYSTAL 0  
nb

The present invention describes a method of inducing and/or sustaining an immunological cytotoxic T lymphocyte (CTL) response in a mammal. The method comprises: (a) delivering an antigen to the mammal at a level to induce an immunological CTL response in the mammal; and (b) maintaining the level of the antigen in the mammal's lymphatic system to maintain

AC Y10601; 12-MAY-1999 (first entry)  
 DE HLA Class I motif peptide SEQ ID NO:531  
 PT Cytotoxic T-lymphocyte response; CTL; antigen; lymphatic system;  
 KW immunisation; tumour; infectious disease; immunotherapy; cancer;  
 KW malignant melanoma; viral disease; hepatitis; AIDS.  
 OS Synthetic.  
 OS Homo sapiens.  
 PN WO9902183-A2.  
 PD 21-JAN-1999.  
 PF 10-JULY-1998; U14289.  
 PR 10-JUN-1997; US-9888320.  
 PR 10-JUL-1997; CA-209815.  
 PA (CTL1-) CTL IMMUNOTHERAPIES CORP.  
 PI Kuendig TM, Simard JJJ,  
 DR WPI; 99-120514/10.  
 PT Inducing a cytotoxic T lymphocyte response - by maintaining a level  
 PT of antigen in the lymphatic system of a mammal so as to provide a  
 sustained CTL response, used to treat, e.g. AIDS  
 Disclosure: Page 49; 199pp; English.  
 PS The present invention describes a method of inducing and/or sustaining  
 an immunological cytotoxic T lymphocyte (CTL) response in a mammal. The  
 method comprises: (a) delivering an antigen to the mammal at a level to  
 induce an immunological CTL response in the mammal, and (b) maintaining  
 the level of the antigen in the mammal's lymphatic system to maintain  
 the immunologic CTL response. The method can be used for the delivery of  
 e.g., a differentiation antigen, a tumour specific multilneage antigen,  
 an embryonic antigen, an oncogene antigen, a mutated tumour-suppressor  
 gene antigen, or a viral antigen. They can be used for the treatment of  
 disease such as cancer, e.g. malignant melanoma or infectious disease,  
 e.g. viral disease such as hepatitis or AIDS. Sustained antigen delivery  
 to the lymphatic system provides for potent CTL stimulation that takes  
 place in the milieu of the lymphoid organ, and it sustains stimulation  
 that is necessary to keep CTL active, cytotoxic and recirculating  
 through the body. Y10071 to Y10639 represent examples of peptide  
 CC antigens given in the present invention.  
 Sequence 9 AA;

Query Match 100.0%; Score 56; DB 1; Length 9;  
 Best Local Similarity 100.0%; Pred. No. 1.03e+01;  
 Matches 9; Conservative 0; Gaps 0;  
 Indels 0; Mismatches 0; Sequence 9 AA;

Db 1 AAGIGILTV 9  
 Qy 1 AAGIGILTV 9  
 Query Match 6 standard; peptide; 9 AA.  
 ID W98938; 06-MAY-1999 (first entry)  
 DE Human leukocyte antigen A2 molecule binding peptide SEQ ID NO:2.  
 KW Human leukocyte antigen; HLA; HLA-A2 binding Peptide; T cell;  
 KW cytolytic T cell; CTL.  
 OS Synthetic.  
 OS Homo sapiens.  
 PN WO958951-A1.  
 PD 30-DEC-1998.  
 PF 18-JUN-1998; U12879.  
 PR 16-APR-1998; US-061388.  
 PR 23-JUN-1997; US-8800963.

PA (LUDWIG INST CANCER RES.)  
 PI Cerottini J, Romero P, Valmori D;  
 DR WPI; 99-105609/09.  
 PT New decamer peptides which bind to HLA molecules - useful to  
 identify HLA-A2 positive molecules and provoke T cells  
 Claim 17: Page 9; 45pp; English.

The present invention describes peptides which bind to an HLA-A2  
 molecule and have Val at the carboxy terminus, and either: (a) Ala, Tyr  
 or Phe at the amino terminus, and Ala at position 2 (P1); or (b) Glu at  
 the amino terminus, and Ala, Leu, or Met at positions 2 and 3, with the  
 proviso that Ala is not at both positions (P2). The present sequence  
 represents an HLA-A2 binding peptide. The peptides of the present

CC invention are used to identify HLA-A2 positive cells, provoke T cells,  
 CC and determine the presence of particular T cells including cytolytic  
 CC T cells (CTLs). They provide a better target than the prior art  
 CC CTL-stimulating peptide.  
 Sequence 9 AA;

Query	Match	Score	DB	Length	Pred. No.	Mismatches	Indels	Gaps
Hom sapiens.	Best Local Similarity 100.0%; Matches 9; Conservative 0;	56	1	9	1.03e+01	0	0	0
W39430	RESULT 7	100.0%	ID	standard; peptide; 9 AA.				
W39430	ID		AC	W39430				
W39430	AC		DT	11-JUN-1998 (first entry)				
W39430	DT		DE	Human immunogenic T cell epitope 1.				
W39430	DE		KW	T cell epitope; immune response; human leukocyte antigen; HLA Class I;				
W39430	KW		KW	vaccine; immunogenic; major histocompatibility complex; MHC; B cell; disease; anti-tumour; anti-viral.				
W39430	KW		OS	Homo sapiens.				
W39430	OS		PN	WO9741440-A1.				
W39430	PN		PD	06-NOV-1997.				
W39430	PD		PF	28-APR-1997; NL0229.				
W39430	PF		PR	23-DEC-1996; EP-203670.				
W39430	PR		PR	26-APR-1996; EP-201145.				
W39430	PR		PA	(YLE-) RIJKSUNIV LEIDEN.				
W39430	PA		PA	(SCIS-) SCI SEED CAPITAL INVESTMENTS BV.				
W39430	PA		P1	Kast WM, Meilef COM, Offringa R, Toes REM, Van Der Burg SH;				
W39430	P1		DR	WPI; 97-549891/50.				
W39430	DR		PT	Method of selecting T cell peptide epitope(s) - by measuring the stability of HLA class I-peptide complexes on intact B cells				
W39430	PT		PS	Disclosure; Page 6; 10pp; English.				
W39430	PS		CC	Peptides W39430-W39734 are used in a novel method for the selection of immunogenic T-cell peptide epitopes present in polypeptides.				
W39430	CC		CC	Peptides W39430 and W39431 are derived from MART-1.				
W39430	CC		CC	The identification of peptide sequences capable of binding to an HLA (human leukocyte antigen) class I molecule and measuring the binding of this epitope peptide to the HLA class I peptide. The stability of binding of the peptide and MHC (major histocompatibility complex) class I				
W39430	CC		CC	CC molecule is measured on intact human B cells carrying the MHC molecule at their cell surfaces. The method can be used to select peptide epitopes for generating vaccines against a disease associated with the peptide, e.g. cancers or AIDS. The peptide epitopes are especially T-cell peptide epitopes with strong anti-tumour and anti-viral immune responses.				
W39430	CC		SQ	Sequence 9 AA;				
W39430	SQ		Query Match 100.0%; Score 56; DB 1; Length 9;					
W39430	Query Match 100.0%; Score 56; DB 1; Length 9;		Best Local Similarity 100.0%; Pred. No. 1.03e+01;					
W39430	Best Local Similarity 100.0%; Pred. No. 1.03e+01;		Matches 9; Conservative 0;					
W39430	Matches 9; Conservative 0;		Gaps 0;					
W39430	Gaps 0;		Indels 0;					
W39430	Indels 0;		Mismatches 0;					
W39430	Mismatches 0;		Sequence 9 AA;					
W39430	Sequence 9 AA;		RESULT 8					
W39430	RESULT 8		ID	WO7379				
W39430	ID		AC	W39430				
W39430	AC		DT	28-JUL-1997 (first entry)				
W39430	DT		DE	MART-1 epitope recognised by melanoma specific T cell receptor.				
W39430	DE		KW	T cell; receptor; lymphocyte; alpha; beta chain; V; variable;				
W39430	KW		J	Joining; D; diversity; gene segment; probe; detection;				
W39430	J		KW	KW recombination; melanoma; cancer; neoplasia; tumour; diagnosis;				
W39430	KW		KW	KW Melanoma Antigen Recognised by T lymphocyte.				
W39430	KW		OS	OS Homo sapiens.				
W39430	OS		PN	WO9630516-A1.				
W39430	PN		PD	03-OCT-1996.				

PF	27-MAR-1996; U04143.	Query Match Score 56; DB 1; Length 9;
PR	27-MAR-1995; US-411098.	Best Local Similarity 100.0%; Pred. No. 1.0e+01;
PA	(USSH ) US DEPT HEALTH & HUMAN SERVICES.	Mismatches 0; Indels 0; Gaps 0;
Hwu P,	Nishimura M,	Rosenberg SA;
DR	WPI: 96-485449/48.	
PT	T cell receptor/alpha and/or beta chains, and related nucleic acids	
PT	useful in pharmaceutical compns. to prevent or treat cancer,	
PT	partic. lung, melanoma, ovarian, colon, brain or kidney tumours	
PS	Example 3; Page 11; 125pp; English	
CC	W07381 are MART-1 epitopes, M9-1, M9-2, M10-3 and M10-4	
CC	respectively, that are recognised by melanoma specific T lymphocyte	
CC	receptors (TCRs). Melanoma-specific TCRs comprising an alpha and	
CC	beta chain were made. Nucleic acids from either of these chains can be	
CC	used as probes for the detection of expression of rearranged genes	
CC	encoding tumour-associated antigens. The nucleic acids may also be used	
CC	to create transgenic animals, useful as biological models to study cancer	
CC	and evaluate diagnostic and therapeutic methods for the treatment of	
CC	cancers, particularly melanomas. Antibodies (Abs) may be raised against	
CC	alpha and beta chain polypeptides and used to detect native or denatured	
CC	TCRs and/or alterations in expression levels of T cells carrying	
CC	melanoma-specific TCRs. Abs can also purify and enrich T cells carrying	
CC	the above receptors, which can then be administered therapeutically to	
CC	mammals. Anti-idiotypic antibodies can be used to assess the level of a	
CC	specific T cell carrying these receptors in a mammal being treated using	
CC	these methods. Host cells and vectors carrying nucleic acid encoding	
CC	a TCR (or individual alpha or beta chain fragment) are useful in	
CC	pharmaceutical compositions to prevent or treat cancer in a mammal, e.g.	
CC	lung, melanoma, ovarian, colon, brain or kidney tumours.	
SQ	Sequence 9 AA;	
Query Match Score 56; DB 1; Length 9;		
Best Local Similarity 100.0%; Pred. No. 1.0e+01;		
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
Db	1 AAGIGILTV 9 	
Qy	1 AAGIGILTV 9	
RESULT 9		
ID	W42523 standard; peptide; 9 AA.	
AC	W42523;	
DT	22-JUN-1998 (first entry)	
DE	Melan A/MART epitope (residues 27-35).	
KW	Metastatic melanoma; peptide analogue; vaccine; cancer; diagnosis;	
KW	antigen; CTL; immunogenic; viral disease; gp 100; Melan A/MART-1.	
KW	Synthetic.	
OS	Homo sapiens.	
PN	W09802538-A1.	
PD	22-JAN-1998; E03712.	
PF	08-JUL-1997; EP-201945.	
PA	(ALKU ) AKZO NOBEL NV.	
PI	Adema GJ, Fijgdr CG;	
PI	WPI: 98-110586/10.	
PT	Melanoma associated peptide analogues - useful in vaccines against	
PT	melanoma	
PS	Example 1; Page 28; 47pp; English.	
CC	This sequence is shown in the specification. The invention relates to	
CC	peptides, which are immunogenic with lymphocytes directed against	
CC	metastatic melanomas. They are characterised in that they comprise at	
CC	least a part of the following sequence, where the amino acid at position	
CC	2 or 8 is substituted: Lys-Thr-Trp-Gly-Gln-Tyr-Trp-Gln-Val. Vaccines	
CC	comprising the peptide, an epitope of the peptide, nucleotide sequence	
CC	encoding the peptide, or an antigen presenting cell preloaded with the	
CC	peptide or antibody as above, are useful for cancer, particularly	
CC	melanoma, treatment. The peptides can also be used to generate antigen	
CC	reactive tumour infiltrating lymphocytes, which can also be used in	
CC	vaccines. The peptides can be exploited to elicit native epitope-reactive	
CC	CTL. Usage of the peptides with improved immunogenicity may contribute	
CC	to the development of CRL-epitope based vaccines in viral disease and	
CC	cancer.	
SQ	Sequence 9 AA;	
Query Match Score 56; DB 1; Length 9;		
Best Local Similarity 100.0%; Pred. No. 1.0e+01;		
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
Db	1 AAGIGILTV 9 	
Qy	1 AAGIGILTV 9	
RESULT 11		
ID	W35512 standard; peptide; 9 AA.	
AC	W35512;	
DT	22-APR-1998 (first entry)	
DE	MART-1/Melan-A protein peptide SEQ ID NO:44 from W09738011.	
KW	T-cell stimulatory peptide; immunogen; non-dendritic carrier; tumour;	
KW	scaffold; inhibition; metastasis; wound healing; solid phase.	
OS	Unidentified.	
PN	W09738011-A1.	
PD	16-OCT-1997.	
PF	03-APR-1997; D00146.	
PA	(PEPR ) PEPPERSEARCH AS	
PA	Heegaard PMH, Jakobsen PH;	
DR	WPI: 97-512645/47.	
PT	Non-dendritic peptide carrier linked to a solid phase - useful as a	
PT	diagnostic agent and as a scaffold for production of chemical	
PT	derivatives	
PS	Example 26; Page 146; 262pp; English.	
CC	A non-dendritic peptide carrier (A) has been developed which is coupled	
CC	through a linker to a solid phase, forming a complex of (A)-solid phase.	
CC	where (A) comprises 10-50 amino acids capable of forming a secondary	
CC	structure in a benign buffer after liberation from the solid phase, and	

further the (A)-solid phase complex comprises an immunogenic substance and/or an immune mediator coupled on (A). The present sequence represents a peptide used in an example from the present invention. An (A)-solid phase complex can be used as a scaffold for the production of chemical derivatives, characterised by covalently attaching molecules at attachment points. Alternatively (A) is used as a scaffold-peptide for the incorporation into an Immunostimulating Complex (Iscom) resulting an (A)-iscom complex which is used for the chemical coupling of antigenic substances in an aqueous solution by conjugation. (A) derivatised with one or more peptides having fibronectin-, laminin- or vitronectin-like binding activities can be used for the promotion of cell-attachment to plastic surfaces, in particular to inhibit tumour growth and metastasis, and for promotion of wound healing. Also a derivatised (A) can be used for the selection of specifically-binding aptamers or as a diagnostic agent. Such diagnostic-(A) molecules could be used to detect molecules derived from or indicative of pregnancy or of a disease, such as an infectious, autoimmune or cancerous disease.

Sequence 9 AA;

Query Match Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 1.03e+01;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

SQ

Db 1 AGIGILTV 9  
| | | | | | | | |  
Qy 1 AGIGILTV 9

RESULT 12  
ID W54602 standard; peptide; 9 AA.  
AC W54602;  
DT 25-SEP-1998 (first entry)  
DE Peptide 1 from Melan-A/Mart-1.  
KW Mannose; antigen; antigen-presenting cell; mannosylated peptide; T cell; vaccine; treatment.  
KW Synthetic.  
OS  
PN W09813378-A1.  
PD 02-APR-1998.  
PF 25-SEP-1997; NL0536.  
PR 26-SEP-1996; EP-202701.  
PA (UYLE-) RIJKSUNIV LEIDEN.  
DR Drijfhout JW, Konig F;  
WPI: 98-230631/20.

PT Increasing uptake and presentation of antigen(s) - by adding mannose residue(s) to antigen for increasing T cell response, useful in, e.g. vaccines against viral infection(s)  
PS Disclosure; Page 24; 47pp; English.  
CC The peptides W54559-W54809 are examples of peptides to which at least 1 (preferably 2) mannose can be attached to increase their uptake as antigens by antigen presenting cells. Uptake of agonist mannosylated peptides will increase the T cell response, whereas uptake of antagonist peptides blocks the T cell response. Blocking binding of immunogenic autoantigens can be used in treatment of type I diabetes, rheumatoid arthritis, graft rejection etc., also to induce T-cell non-responsiveness. Vaccines containing mannosylated antigen are used to prevent or treat infections by, e.g. bacteria, viruses, fungi, helminths and parasites.  
Sequence 9 AA;

Query Match Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 1.03e+01;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

SQ

Db 1 AGIGILTV 9  
| | | | | | | | |  
Qy 1 AGIGILTV 9

RESULT 13  
ID W77123 standard; peptide; 9 AA.  
AC W77123;  
DT 16-NOV-1998 (first entry)  
DE MART-1/MelanA synthetic peptide epitope 1.

KW Tyrosinase; tyrosinase cytotoxic lymphocyte response;  
cytotoxic T lymphocyte; cysteine-depleted; melanoma.  
OS Synthetic.

PN W09833810-A2.  
PD 06-AUG-1998.  
PF 29-JAN-1998; U01592.  
PR 30-JAN-1997; US037781.

PA (UVVI-) UNIV VIRGINIA PATENT FOUND.  
PI Engelhard VH, Hunt DF, Kittleson D, Slingluff CL;  
WPI; 98-437388/37.  
DR - comprises disease specific cytotoxic T lymphocyte epitope used to elicit melanoma specific CTL response.  
PT Disclosure; Page 27; 93pp; English.  
PS The peptide epitope W77119-W77138 were created for human tumour-specific cytotoxic T lymphocyte response. These peptides are cysteine-depleted mutants of a native disease-specific CTL epitope. The cysteine-depleted CTL epitopes elicit a stronger or more specific CTL response than the native epitope. The epitopes can be used in a disease-specific immunogen to protect a mammal against disease in particular melanomas.  
CC The peptides may also be used to screen a sample for the presence of an antigen with the same epitope, or with a different cross-reactive epitope.

CC Sequence 9 AA;

Query Match Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 1.03e+01;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

SQ

Db 1 AAGIGILTV 9  
| | | | | | | | |  
Qy 1 AAGIGILTV 9

RESULT 14  
ID W68380 standard; peptide; 9 AA.  
AC W68380;  
DT 14-OCT-1998 (first entry)  
DE Human MART1/MELAN-A peptide binds HLA-A2.  
KW Antigen; major histocompatibility complex; MHC; lymphocyte; detection;  
KW immobilisation; cytotoxic T-cell; tumour; leukaemia; lymphoma;  
KW viral infection.  
OS Synthetic.  
OS Homo sapiens.  
PN W09744667-A2.  
PD 27-NOV-1997.  
PF 21-MAY-1997; F00892.  
PR 21-MAY-1996; US-651925.  
PA (INRM ) INSERM INST NAT SANTE & RECH MEDICALE.  
PA (INSP ) INST PASTEUR.  
PI Abastado J, Kourilsky P, Langlade-Demoyen P, Lone Y;  
DR WPI; 98-018653/02.  
PT Detection, purification and elimination of antigen-specific T cells for immuno-therapy of cancers and viral infection.  
PT Cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2). A similar method is used to isolate, purify or eliminate Ag-specific T-cells or to produce Ag-specific cytotoxic T-cells (cTcC). The method is also used to detect and quantify tumour-specific T-cells and to generate CTC for specific killing of tumour cells (solid tumours, leukaemia or lymphoma) by injection into a human or animal, but also for treating viral infections.  
CC Onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

CC Peptides W68301-W68384 are examples of antigens (Ag) which can be loaded onto recombinantly produced major histocompatibility complex (MHC) molecules in a method of detecting antigen specific lymphocytes. The MHC-antigen complex is then immobilised on a solid support and a sample containing cells recognising the MHC-Ag complex may be isolated. This peptide is derived from the human MART1/MELAN-A protein and binds the human leukocyte antigen A2 (HLA-A2).

Query Match Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 1.03e+01;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AAGIGILTV 9  
 AC |||||||  
 QY 1 AAGIGILTV 9

RESULT 15  
 ID Y01750; standard; Peptide; 10 AA.

AC Y01750;  
 DT 25-JUN-1999 (first entry)  
 DE Exemplary antigenic peptide derived from Melan-A(MART-1).  
 KW MAGE-3; tumour associated gene; human leucocyte antigen Class II;  
 autologous CD4+ cell; MAGE-3 related disease; cancer; melanoma;  
 osteosarcoma; leukemia; carcinoma.  
 OS Homo sapiens.  
 PN W09914326-A1.  
 PD 25-MAR-1999.  
 PF 04-SEP-1998; U18601.  
 PR 12-SEP-1997; US-928615.

PA (LUDWIG INST CANCER RES.

PA (UYVR) UNIV VRIJE BRUSSEL.

PI Boon-Faïleur T, Chaux P, Corthals J, Heirman C,

PJ Luiten R, Stroobant V, Thielemans K, Van Der Bruggen P;

DR WPI: 99-244031/20.

PT Isolated peptides that bind to human leucocyte antigen class II

PS Disclosure: Page 29; 88PP; English.

CC The present sequence represents an exemplary tumour associated peptide CC antigen. The specification describes a MAGE-3 tumour associated gene. CC Peptides (Y01721-25) that bind human leucocyte antigen (HLA) Class II CC molecules can be derived from the MAGE-3 protein. These peptides and CC autologous CD4+ cells that bind to a complex of MAGE-3 peptide CC and HLA Class II are used to treat MAGE-3 related diseases, CC particularly cancers (e.g. melanoma, osteosarcoma, leukemia and CC various forms of carcinoma). The peptides are also used to produce CC specific antibodies. Detection of the peptides, e.g. in binding CC assays, particularly with antibodies, is used for diagnosis of such CC diseases.

Sequence 10 AA:

Query Match 100.0%; Score 56; DB 1; Length 10;  
 Best Local Similarity 100.0%; Pred. No. 1.03e+01;  
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 2 AAGIGILTV 10  
 AC |||||||  
 QY 1 AAGIGILTV 9

Search completed: Fri May 5 21:59:14 2000  
 Job time : 35 secs.



KW ~autologous CD4+ cell; MAGE-3 related disease; cancer; melanoma;  
 KW osteosarcoma; leukemia; carcinoma.  
 OS Homo sapiens.

PN - WO9914226-A1.

PD 25-MAR-1999.

PF #4-SEP-1998; 18801.

PR 12-SEP-1997; US-528615.

{ LUDWIG INST CANCER RES.  
 PA (LUDWIG) LUDWIG INST CANCER RES.  
 PA (UVR) UNIV VRIJE BRUSSEL.

PI Boon-Falleur T, Chaux P, Corthals J, Heirman C,  
 PI Luijten R, Stroobant V, Thielemans K, Van Der Bruggen P;

DR WPI; 99/244031/20.

PT Isolated peptides that bind to human leucocyte antigen class II  
 PR molecules

PS Disclosure; Page 29; 88pp; English.

CC The present sequence represents an exemplary tumour associated peptide  
 CC antigen. The specification describes a MAGE-3 tumour associated gene.  

CC Peptides (Y0721-25) that bind human leucocyte antigen (HLA) Class II  
 CC molecules can be derived from the MAGE-3 protein. These peptides and  
 CC autologous CD4+ cells that bind to a complex of MAGE-3 peptide  
 CC and HLA Class II, are used to treat MAGE-3 related diseases,  
 CC particularly cancers (e.g. melanoma, osteosarcoma, leukemia and  
 CC various forms of carcinoma). The peptides are also used to produce  
 CC specific antibodies. Detection of the peptides, e.g. in binding  
 CC assays, particularly with antibodies, is used for diagnosis of such  
 CC diseases.

CC Sequence 10 AA;

SQ Score 62; DB 1; Length 10;  
 Best Local Similarity 100.0%;  
 Matches 10; Conservative 0; Gaps 0;

Db 1 EAAGIGILTV 10

Qy 1 EAAGIGILTV 10

RESULT 4  
 ID W39447 standard; peptide; 10 AA.  
 AC W39447;  
 DT 11-JUN-1998 (first entry)  
 DE Human HLA-A\*0201 immunogenic peptide 10-mer.  
 KW T cell epitope; immune response; human leukocyte antigen; HLA Class I;  
 KW vaccine; immunogenic; major histocompatibility complex; MHC; B cell;  
 KW disease; anti-tumour; anti-viral.  
 OS Synthetic.  
 OS Homo sapiens.  
 PN WO9741440-A1.  
 PD 06-NOV-1997.  
 PF 28-APR-1997; NL0229.

PR 23-DEC-1996; EP-203670.  
 PR 26-APR-1996; EP-201145.  
 PA (UYLE-) RIJKSUNIV LEIDEN.  
 PA (SCIS-) SCI SEED CAPITAL INVESTMENTS BV.  
 PI Kast WM, Melief CJJM, Offringa R, Toes REM, Van Der Burg SH;  
 DR WPI; 97-51981/50.

PT Method of selecting T cell peptide epitope(s) - by measuring the  
 PT stability of HLA class I peptide complexes on intact B cells  
 PS Example 3; Page 29; 109pp; English.

CC Peptides W39430-W39734 are used in a novel method for the selection of  
 CC immunogenic T-cell peptide epitopes present in polypeptide antigens. The  
 CC method involves the identification of peptide sequences capable of  
 CC binding to an HLA (human leukocyte antigen) class I molecule and  
 CC measuring the binding of this peptide to the HLA class I peptide.  
 CC The stability of binding of the peptide and MHC (major histocompatibility  
 CC complex) class I molecule is measured on intact human B cells carrying  
 CC the MHC molecule at their cell surfaces. The method can be used to select  
 CC peptide epitopes for generating vaccines against a disease associated  
 CC with the polypeptide, e.g. cancers or AIDS. The peptide epitopes are  
 CC especially T-cell peptide epitopes with strong anti-tumour and anti-viral  
 CC immune responses. Peptide W39447 is an immunodominant peptide-epitope  
 CC presented by HLA-A\*0201-positive melanoma cells and displays considerable  
 CC binding to HLA-A\*0201 in assays.

SQ Sequence 10 AA;

Query Match Score 62; DB 1; Length 10;  
 Best Local Similarity 100.0%;  
 Matches 10; Conservative 0; Gaps 0;

Db 1 EAAGIGILTV 10

Qy 1 EAAGIGILTV 10

RESULT 5  
 ID W07380 standard; Peptide; 10 AA.  
 AC W07380;  
 DT 28-JUL-1997 (first entry)  
 DE MART-1 epitope recognised by melanoma specific T cell receptor.  
 KW T cell; receptor; lymphocyte; alpha; beta chain; V; variable;  
 KW J; joining; D; diversity; gene segment; probe; detection;  
 KW recombination; melanoma; cancer; neoplasia; tumour; diagnosis;  
 KW MART; Melanoma Antigen Recognised by T lymphocyte.  
 OS Homo sapiens.  
 PN WO9630516-A1.  
 PD 03-OCT-1996.

PF	27-MAR-1996; U04143.	Db	1 EAAGIGILTV 10
PA	27-MAR-1995; US-411098.	OY	1 EAAGIGILTV 10
PA	(USSH ) US DEPT HEALTH & HUMAN SERVICES.		
PI	Hwu P, Nishimura M, Rosenberg SA;		
DR	WPI; 96-485449/48.		
PT	T cell receptor alpha and/or beta chains, and related nucleic acids	RESULT	7
PT	- useful in pharmaceutical compns. to prevent or treat cancer,	ID	W54809 standard; peptide; 10 AA.
PT	partic. Lung, melanoma, ovarian, colon, brain or kidney tumours	AC	W54809;
PS	Example 3: Page 11; 125pp; English	DT	26-SEP-1998 (first entry)
CC	W07378-W07381 are MART-1 epitopes M9-1, M9-2, M10-3 and M10-4	DE	Peptide 1 from Mart-1/Melan A.
CC	respectively, that are recognised by melanoma specific T lymphocyte	KW	Mannose; antigen; antigen-presenting cell; mannosylated peptide; T cell
CC	receptors (TCRs). Melanoma-specific TCRs comprising an alpha and	KW	vaccine; treatment.
CC	beta chain were made. Nucleic acids from either of these chains can be	OS	Synthetic.
CC	used as probes for the detection of expression of rearranged genes	PN	WO9813378-A1.
CC	encoding tumour-associated antigens. The nucleic acids may also be used	PD	02-APR-1998
CC	to create transgenic animals useful as biological models to study cancer	PF	25-SEP-1997; NL0536.
CC	and evaluate diagnostic and therapeutic methods for the treatment of	PR	26-SEP-1996; EP-202701.
CC	cancers, particularly melanomas. Antibodies (Abs) may be raised against	PA	(UYLE-) RIJJSUNIV LEIDEN.
CC	alpha and beta chain polypeptides and used to detect native or denatured	PI	Drijfhout JW, Koning F;
CC	TCRs and/or alterations in expression levels of T cells carrying	DR	WPI; 98-230531/20.
CC	melanoma-specific TCRs. Abs can also purify and enrich T cells carrying	PT	Increasing uptake and presentation of antigen(s) - by adding mannose
CC	the above receptors, which can then be administered therapeutically to	PT	residue(s) to antigen for increasing T cell response, useful in,
CC	mammals. Anti-diotype antibodies can be used to assess the level of a	PT	e.g. vaccines against viral infection(s)
CC	specific T cell carrying these receptors in a mammal being treated using	PS	Disclosure: Page 25; 47pp; English.
CC	these methods. Host cells and vectors carrying nucleic acid encoding	CC	The peptides W54559-W54809 are examples of peptides to which at least 1
CC	a TCR (or individual alpha or beta chain fragment) are useful in	CC	(preferably 2) mannose can be attached to increase their uptake as
CC	pharmaceutical compositions to prevent or treat cancer in a mammal, e.g.	CC	antigen-presenting cells. Uptake of agonist mannosylated
CC	lung, melanoma, ovarian, colon, brain or kidney tumours. In a mammal, e.g.	CC	peptides will increase the T cell response, whereas uptake of antagonist
CC	sequence 10 AA;	CC	peptides blocks the T cell response. Blocking binding of immunogenic
SQ	Query Match 100.0%; Score 62; DB 1; Length 10;	CC	autoantigens can be used in treatment of type I diabetes, rheumatoid
Matches 10; Conservative 0; Indels 0; Gaps 0;	Pred. No. 2.79e+00;	CC	arthritis, graft rejection etc., also to induce T-cell non-
Db	1 EAAGIGILTV 10	CC	responsiveness. Vaccines containing mannosylated antigen are used to
QY	1 EAAGIGILTV 10	CC	prevent or treat infections by, e.g. bacteria, viruses, fungi, helminths
Query Match 100.0%; Score 62; DB 1; Length 10;	Sequence 10 AA;	SO	and parasites.
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	Query Match 100.0%; Score 62; DB 1; Length 10;	Query	
Matches 10; Conservative 0; Indels 0; Gaps 0;	Best Local Similarity 100.0%; Pred. No. 2.79e+00;	Match	
Db	1 EAAGIGILTV 10	Matches	10; Conservative 0; Indels 0; Gaps 0;
QY	1 EAAGIGILTV 10	Db	1 EAAGIGILTV 10
Query Match 100.0%; Score 62; DB 1; Length 10;	Best Local Similarity 100.0%; Pred. No. 2.79e+00;	QY	1 EAAGIGILTV 10
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	Query Match 100.0%; Score 62; DB 1; Length 10;	RESULT	8
Matches 10; Conservative 0; Indels 0; Gaps 0;	Best Local Similarity 100.0%; Pred. No. 2.79e+00;	ID	W22039 standard; peptide; 10 AA.
Db	1 EAAGIGILTV 10	AC	W22039;
QY	1 EAAGIGILTV 10	DT	20-FEB-1998 (first entry)
Query Match 100.0%; Score 62; DB 1; Length 10;	Antigenic MART-1 peptide M10-3.	DE	Antigenic MART-1 peptide M10-3.
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	Antigenic peptide; human papillomavirus; MART-1; M10-3; MAGE gene;	KW	Antigenic peptide; human papillomavirus; MART-1; M10-3; MAGE gene;
Matches 10; Conservative 0; Indels 0; Gaps 0;	Human immunodeficiency virus; cancer antigen; tyrosinase; signal protein;	KW	Human immunodeficiency virus; cancer antigen; tyrosinase; signal protein;
Db	1 EAAGIGILTV 10	DE	Human immunodeficiency virus; cancer antigen; tyrosinase; signal protein;
QY	1 EAAGIGILTV 10	PA	(HARD ) HARVARD COLLEGE.
Query Match 100.0%; Score 62; DB 1; Length 10;	Ballard JD, Blanke SR, Collier RJ, Lyszzak EL, Milne JC;	PA	Ballard JD, Blanke SR, Collier RJ, Lyszzak EL, Milne JC;
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	Human sapiens.	OS	Human sapiens.
Matches 10; Conservative 0; Indels 0; Gaps 0;	PN	PN	W09723236-A1.
Db	1 EAAGIGILTV 10	PD	03-JUL-1997; U20463.
QY	1 EAAGIGILTV 10	PF	13-DEC-1996; U20463.
Query Match 100.0%; Score 62; DB 1; Length 10;	Introducing therapeutic proteins, especially antigens, into cells	PT	Introducing therapeutic proteins, especially antigens, into cells
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	- using toxin molecules and/or polycationic handles for delivery	PT	- using toxin molecules and/or polycationic handles for delivery
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PR	07-JUN-1996; US-019275.
QY	1 EAAGIGILTV 10	PR	13-DEC-1995; US-019518.
Query Match 100.0%; Score 62; DB 1; Length 10;	(HARD ) HARVARD COLLEGE.	PA	(HARD ) HARVARD COLLEGE.
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	Ballard JD, Blanke SR, Collier RJ, Lyszzak EL, Milne JC;	PA	Ballard JD, Blanke SR, Collier RJ, Lyszzak EL, Milne JC;
Matches 10; Conservative 0; Indels 0; Gaps 0;	Human sapiens.	OS	Human sapiens.
Db	1 EAAGIGILTV 10	PI	PI
QY	1 EAAGIGILTV 10	DR	Starnbach MN, WPI; 97-350182/32.
Query Match 100.0%; Score 62; DB 1; Length 10;	Introducing therapeutic proteins, especially antigens, into cells	PT	Introducing therapeutic proteins, especially antigens, into cells
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	- using toxin molecules and/or polycationic handles for delivery	PT	- using toxin molecules and/or polycationic handles for delivery
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	(LUDW ) LUDWIG INST CANCER RES.	PS	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	Jager E, Knuth A;	PS	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	WPI; 97-415070/38.	PS	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	Composition containing immunogen and granulocyte macrophage colony stimulating factor; GM-CSF; adjuvant.	PS	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	GM-CSF; adjuvant.	PS	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	PN	PS	CC
Db	1 EAAGIGILTV 10	PD	03-JUL-1997; U20463.
QY	1 EAAGIGILTV 10	PF	13-DEC-1996; U20463.
Query Match 100.0%; Score 62; DB 1; Length 10;	This sequence represents a specifically claimed example of a tumour	PT	Introducing therapeutic proteins, especially antigens, into cells
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	rejection antigen (TRA) which was used with granulocyte	PT	- using toxin molecules and/or polycationic handles for delivery
Matches 10; Conservative 0; Indels 0; Gaps 0;	colony-stimulating factor (GM-CSF) as adjuvant to generate an immune,	PS	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	specifically cytolytic T cell (CTL), response for treatment of cancers	PS	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	or where cell transformation has occurred, e.g. in melanoma or dysplastic	PS	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	nevii. These tumour rejection antigens can also be used diagnostically (if	PS	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	they can induce CTL or antibodies specific for the antigens then this	PS	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	increases, immune response to the tumour rejection antigens.	PS	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	Sequence 10 AA;	PS	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10	PS	CC
QY	1 EAAGIGILTV 10	PS	CC
Query Match 100.0%; Score 62; DB 1; Length 10;	CC	CC	CC
Best Local Similarity 100.0%; Pred. No. 2.79e+00;	CC	CC	CC
Matches 10; Conservative 0; Indels 0; Gaps 0;	CC	CC	CC
Db	1 EAAGIGILTV 10</		

CC compound is linked to either of the delivery molecules by a covalent bond. The B moiety of a toxin enhances delivery of the antigenic compound into a cell. The anthrax toxin system of the invention eliminates the CC problems associated with incorrect folding of lengthy fusion proteins. CC Small cationic fusion peptides substituted for LFN may reduce the possibility of biological activity of the translocated protein. The method is used for the introduction of the CC antigens, e.g. MHC Class I antigens or any other therapeutic protein, CC e.g. toxin molecules, apoptosis-inducing molecules or signalling proteins into the cells.

SQ

Query Match

Best Local Similarity 100.0%

Score 62; DB 1; Length 10;

Pred. No. 2.79e+00;

Matches 10; Conservative 0;

MisMatches 0; Indels 0;

Gaps 0;

PT

CC

AC

ID R84197 standard; Peptide: 10 AA.

AC R84197;

ID 20-APR-1996 (first entry)

DE MART-1 melanoma antigen immunogenic peptide M10-3 derivative.

KW M10-3; melanoma; tumour-associated antigen; melanoma;

KW metastatic melanoma; tumour-associated antigen;

KW immunogenic peptide; diagnosis; prognosis; prophylaxis;

KW therapy; vaccine.

OS Synthetic.

PN WO529193-A2.

PD 02-NOV-1995.

PF 21-APR-1995; US-05063.

PR 22-APR-1994; US-231565.

PT 05-APR-1995; US-417174.

PA (USSH ) US SEC DEPT HEALTH.

PI Kawakami Y, Rosenberg SA;

DR WPI: 95-382963/49.

PT DNA encoding melanoma antigens recognised by T-lymphocytes - also

PT vectors, host cells and antibodies, used to detect, treat and

PT immunise animal against melanoma.

PS Claim 12; Page 122; 184pp; English.

CC Immunogenic peptide M10-3 is a derivative of peptide M9-2 (R84196)

CC which is based on the melanoma antigen (MART-1) (see R84212).

CC M9-2 may be modified to improve immunogenicity (see R84783-R84800)

CC and used in medicaments for the treatment or prevention (by

CC immunogenic peptides) of melanoma. Antibodies against MART-1 and its

CC immunogenic peptides may be used in the detection and isolation of

CC MART-1 from a sample, the detection of which is indicative of a

CC disease state (melanoma or metastatic melanoma).

CC See also R84198.

SQ Sequence 10 AA;

Query Match

Best Local Similarity 100.0%

Score 62; DB 1; Length 10;

Pred. No. 2.79e+00;

Matches 10; Conservative 0;

MisMatches 0; Indels 0;

Gaps 0;

PT

CC

AC

ID R84122 standard; Peptide: 118 AA.

AC R84122;

ID 20-APR-1996 (first entry)

DE MART-1 melanoma antigen.

KW M10-3; melanoma; tumour-associated antigen; melanoma;

KW metastatic melanoma; tumour-associated antigen; immunogen;

KW diagnosis; prognosis; prophylaxis; therapy; vaccine.

OS Mammalian.

FH Key region

FT 27. 47

FT /note= "hydrophobic region"

FT WO529193-A2.

FT PD 02-NOV-1995;

FT PF 21-APR-1995; US-05063.

FT PR 22-APR-1994; US-231565.

FT PR 05-APR-1995; US-417174.

FT PA (USSH ) US SEC DEPT HEALTH.

FT PI Kawakami Y, Rosenberg SA;

FT DR WPI: 95-382963/49.

FT DR N-PSDB; T02714.

FT DNA encoding melanoma antigens recognised by T-lymphocytes - also

FT PT vectors, host cells and antibodies, used to detect, treat and

FT PT immunise animal against melanoma.

FT PS Claim 11; Page 117; 184pp; English.

FT The melanoma antigen (MART-1) is produced by recombinant DNA

FT CC methods, i.e. preferably using a baculovirus vector for expression

FT CC in insect cell cultures. MART-1 protein is a source of immunogenic

FT CC peptides (see R84196 for peptide M9-2) which are optionally modified

FT CC (see R84783-R84800) and used in medicaments for the treatment of

FT CC prevention (by immunization) of melanoma. Antibodies against MART-1

FT CC and its immunogenic peptides may be used in the detection and

FT CC isolation of MART-1 from a sample, the detection of which is

FT CC indicative of a disease state (melanoma or metastatic melanoma).

FT SQ Sequence 118 AA;

Query Match

Best Local Similarity 100.0%

Score 62; DB 1; Length 118;

Pred. No. 2.79e+00;

Matches 10; Conservative 0;

MisMatches 0; Indels 0;

Gaps 0;

PT

CC

AC

ID R84158 standard; Protein: 118 AA.

AC R63158;

ID R63158; (first entry)

AC R63158;

DT 26-MAY-1995 (first entry)

DE Tumour rejection antigen precursor.

KW Isolation; melanoma; cell line; LB-39-MEL; diagnosis; vaccine;

KW therapy.

OS Homo sapiens.

PN WO9421126-A.

PD 29-SEP-1994;

PF 09-MAR-1994; 002487.

PR 18-MAR-1993; US-032978.

PA (LUDWIG INST CANCER RES).

PI Bonn-Failler T, Brichard V, De Plaen E, Traversari C;

PI Van Pel A, Wolfel T;

DR WPI: 94-316544/39.

DR N-PSDB; Q76370.

PT Nucleic acid coding for a tumour rejection antigen precursor - is

PT used for developing prdt. for diagnosis or treatment of expression

PT related disorders, partic. melanoma

PS Claim 5; Page 14; 264pp; English.

CC This sequence represents the tumour rejection antigen presented by HLA-A2 molecules.

CC The tumour rejection antigen is not related to tyrosinase. The cDNA

CC encoding this sequence was isolated from the melanoma cell line,

CC LB-39-MEL. The tumour rejection antigen may be used for diagnosis or

CC in vaccines or for therapy of disorders characterised by the expression

CC of the tumour rejection antigen precursor, particularly melanoma.

SQ Sequence 118 AA;

Query Match

Best Local Similarity 100.0%

Score 62; DB 1; Length 118;

Pred. No. 2.79e+00;

Matches 10; Conservative 0;

MisMatches 0; Indels 0;

Gaps 0;

Db 26 EAAGIGILTV 35

|||||||

QY 1 EAAGIGILTV 10  
 RESULT 12  
 ID W8134 standard; peptide; 118 AA.  
 AC W8134;  
 DT 04-FEB-1999 (first entry)  
 DE Human; tumour rejection antigen precursor; human leukocyte antigen;  
 KW TRAP; HLA; cancer; melanoma.  
 OS Homo sapiens.  
 FH Key  
 FT Location/Qualifiers  
 Misc\_difference 2  
 /note= "encoded by CGA"  
 FR Misc\_difference 17  
 /note= "encoded by GAC"  
 FT PNT 05537476-A.  
 PD 17-NOV-1998.  
 PF 16-JAN-1998; 007966.  
 PR 03-MAR-1995; US-398409.  
 PA (LUDWIG INST CANCER RES.  
 PI Boon-Falleur T, Brichard V, De Plaein E, Traversari C,  
 Van Pel A, Woelfelt;  
 WPI; 99-043967/04.  
 DR N-BSD; V70150.  
 PR Use of a tumour rejection antigen precursor - as a marker for  
 PT diagnosing a disorder characterised by expression of a tumour  
 PR rejection antigen precursor which is not tyrosinase  
 PS Claim 1; Column 7, 9; 11pp; English.  
 CC A method has been developed for the diagnosis of a disorder which is  
 characterised by the expression of a tumour rejection antigen precursor  
 (TRAP) which is not tyrosinase, and which is processed to a TRA which  
 forms a complex with an HLA-A2 molecule. The present sequence represents  
 the TRAP for use in the present invention. The method comprises  
 contacting a sample from subject with an agent specific for the  
 complex and determining the interaction between the complex and the  
 agent as a determination of the disorder. TRAP can be used for the  
 diagnosis and treatment of disorders characterised by the expression  
 of the TRAP molecules such as cancers, particularly melanoma.  
 Sequence 118 AA;

Query Match Score 62; DB 1; Length 118;  
 Best Local Similarity 100.0%; Pred. No. 2.79e+00;  
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 SQ

Db 26 EAAGIGILTV 35  
 Qy 1 EAAGIGILTV 10

RESULT 13  
 ID W98336 standard; peptide; 10 AA.  
 AC W98336;  
 DT 06-MAY-1999 (first entry)  
 DE Human leukocyte antigen A2 molecule binding peptide SEQ ID NO:24.  
 KW Human leukocyte antigen; HLA; HLA-A2 binding peptide; T cell;  
 cytolytic T cell; CTL.  
 OS Homo sapiens.  
 PN W09858951-A1.  
 PD 30-DEC-1998.  
 PR 18-JUN-1998; U12879.  
 PR 16-APR-1998; US-061388.  
 PR 23-JUN-1997; US-880963.  
 PA (LUDWIG INST CANCER RES.  
 PI Cerottini J, Romero P, Vallmori D;  
 WPI; 99-105609/09.  
 PR New decamer peptides which bind to HLA molecules - useful to  
 identify HLA-A2 positive cells and provoke T cells.  
 Claim 13; Page 21; 45pp; English.  
 The present invention describes peptides which bind to an HLA-A2  
 molecule and have Val at the carboxy terminus, and either: (a) Ala, Tyr

CC or Phe at the amino terminus, and Ala at position 2 (P1); or (b) Glu at  
 the amino terminus, and Ala, Leu, or Met at positions 2 and 3, with the  
 proviso that Ala is not at both positions (P2). The present sequence  
 represents an HLA-A2 binding peptide. The peptides of the present  
 invention are used to identify HLA-A2 positive cells, provoke T cells,  
 and determine the presence of particular T cells including cytolytic  
 T cells (CTLs). They provide a better target than the prior art  
 CTL-stimulating peptide.

SQ Sequence 10 AA;

Query Match Score 59; DB 1; Length 10;  
 Best Local Similarity 90.0%; Pred. No. 6.17e+00;  
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Db 1 EAAGIGILAV 10  
 Qy 1 EAAGIGILTV 10

RESULT 14  
 ID Y10567 standard; peptide; 9 AA.  
 AC Y10567;  
 DT 12-MAY-1999 (first entry)  
 DE HLA Class I motif peptide SEQ ID NO:497.  
 KW Cytotoxic T-lymphocyte response; CTL; antigen; lymphatic system;  
 immunisation; tumour; infectious disease; immunotherapy; cancer;  
 malignant melanoma; viral disease; hepatitis; AIDS.  
 OS Synthetic.

OS Homo sapiens.  
 PN W090211-A2.  
 PD 21-JAN-1999.  
 PR 10-DEC-1997; US-988320.  
 PR 10-JUL-1997; CA-209815.  
 PA (CTLI-) CTL IMMUNOTHERAPIES CORP.  
 PI Kuendig TM, Simard JJL;  
 DR WPI; 99-120514/10.  
 PR Inducing a cytotoxic T lymphocyte response - by maintaining a level  
 PT of antigen in the lymphatic system of a mammal so as to provide a  
 sustained CTL response, used to treat, e.g. AIDS.  
 PS Disclosure; Page 47; 199pp; English.

CC The present invention describes a method of inducing and/or sustaining  
 an immunological CTL response in a mammal. The  
 method comprises: (a) delivering an antigen to the mammal at a level to  
 induce an immunological CTL response in the mammal's lymphatic system to maintain  
 the level of the antigen in the mammal's lymphatic system to maintain  
 the immunologic CTL response. The method can be used for the delivery of  
 e.g. a differentiated antigen, a tumour-specific multileague antigen,  
 an embryonic antigen, an oncogene antigen, a mutated tumour-suppressor  
 gene antigen, or a viral antigen. They can be used for the treatment of  
 CC disease such as cancer, e.g. malignant melanoma or infectious disease,  
 CC e.g. viral disease such as hepatitis or AIDS. Sustained antigen delivery  
 CC to the lymphatic system provides for potent CTL stimulation that takes  
 place in the milieu of the lymphoid organ, and it sustains stimulation  
 CC that is necessary to keep CTL active, cytotoxic and recirculating  
 CC through the body. Y10071 to Y10639 represent examples of peptide  
 CC sequences given in the present invention.  
 SQ Sequence 9 AA;

Query Match Score 56; DB 1; Length 9;  
 Best Local Similarity 100.0%; Pred. No. 1.30e+01;  
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 EAAGIGILTV 9  
 Qy 2 EAAGIGILTV 10

RESULT 15  
 ID Y00713 standard; peptide; 9 AA.  
 AC Y00713;  
 DT 12-MAY-1999 (first entry)  
 DE Tumour antigen booster peptide Melan-AMAP-1 HLA-A2 #2.

KW Tumour antigen; booster peptide; immune response modulation; allergy;  
 KW immune response enhancer; tumour cell; tumour rejection antigen;  
 KW leukocyte antigen presenting molecule; autoimmune disease;  
 KW allograft rejection.  
 OS Homo sapiens.  
 PN WO959956-A2.  
 PD 30-DEC-1998.  
 PF 19-JUN-1998; U12894.  
 PR 23-JUN-1997; US-888979.  
 PA (LUDWIG) LUDWIG INST CANCER RES.  
 PI Boon-Falleur T, Uyttenhove C, Warnier G;  
 DR WPI; 99-105612/09.  
 PT Immunization methods using viruses expressing antigen for priming  
 and booster immunizations - useful for modulating immune responses  
 against antigen, e.g. enhancing immune response against tumour cells  
 PT expressing tumour rejection antigens  
 PS Disclosure; Page 10; 33BP; English.  
 CC This sequence represents a tumour antigen booster peptide that can be  
 CC used in the method of the invention. The method is for modulating an  
 CC immune response in a mammal against an antigen, and comprises:  
 CC (A) inducing an immune response by: (1) administering a virus containing  
 CC a nucleic acid molecule encoding the antigen or its precursor to generate  
 CC an immune response; and (ii) administering at least one booster dose  
 CC comprising a peptide including the antigen, in an adjuvant, in a combined  
 CC amount effective to enhance the initial immune response; or  
 CC (B) reducing an immune response as defined for (A) but using a  
 CC non adjuvant with the peptide which includes the antigen, in an amount  
 CC effective to reduce the initial immune response. Method (A) is used to  
 CC enhance the immune response against tumour cells expressing tumour  
 CC rejection antigens, and against pathogens in subjects having human  
 CC leukocyte antigen presenting molecules. Method (B) is used to reduce the  
 CC immune response in allergy, autoimmune disease, and allograft rejection.  
 CC Method (A) provides an immunisation method which, unlike prior art, is  
 CC not limited by the host immune response against viral vectors.  
 SQ Sequence 9 AA;

Query Match 90.3%; Score 56; DB 1; Length 9;  
 Best Local Similarity 100.0%; Pred. No. 1.36e+01;  
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Db 1 AAGTGTLY 9  
 Qy ||||||| 2 AAGTGTLY 10

Search completed: Fri May 5 22:07:42 2000  
 Job time : 33 secs.

The logo consists of the letters "TM" in a stylized, blocky font. The "T" is a vertical rectangle with a horizontal bar extending from its top right corner. The "M" is formed by two diagonal lines meeting at their midpoints, creating a V-shape.

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on: Fri May 5 22:10:16 2000; MasPar time 7.35 Seconds  
Serial output not generated.  
94.341 Million cell updates/sec

>PS-09-267-439-17

Scripture: 1 Peter 1:10 from US0926/439.pep  
Perfect Score: 62  
Accuracy: 100

scoring table: PAM 150

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archaea:      25878 seqs, 6934122 residues
st-processing: Minimum Match 08
                  Listing first 45 summaries
database:      sptrembl12
1:sp-archaea 2:sp_bacteria 3:sp_fungi 4:sp_human
5:sp_invertebrate 6:sp_mammal 7:sp_mhc 8:sp_organelle
9:sp_phage 10:sp_plant 11:sp_rodent 12:sp_unclassified
13:sp_virus 14:sp_virion
15:sp_vertebrate

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statistics: Mean 22.226; Variance 25.879; scale 0.059  
 Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match Length DB ID			Description	Pred. No.
		Match	Length	DB		
1	53	85.5	766	10	Q23161 RECEPTOR KINASE-LIKE P	7.4e-01
2	48	77.4	808	10	Q9ZT37 PUTATIVE GLUTAMATE REC	9.67e+00
3	47	75.8	165	2	P71810 HYPOTHETICAL 18.2 KD P	1.58e+01
4	47	75.8	250	2	Q31597 YJBA PROTEIN.	1.58e+01
5	47	75.8	478	2	Q87090 GLYCOPROTEIN GIII.	1.58e+01
6	47	75.8	479	14	Q87089 GLYCOPROTEIN GIII.	1.58e+01
7	47	75.8	479	14	Q87091 GLYCOPROTEIN GIII.	1.58e+01
8	47	75.8	773	10	Q22178 PUTATIVE RECEPTOR PROT	1.58e+01
9	47	75.8	2219	5	Q23388 ZK1067.2 PROTEIN.	1.58e+01
10	46	74.2	848	5	O18139 T26H2.7 PROTEIN.	2.57e+01
11	46	74.2	1347	2	O30426 XYLANASE.	2.57e+01
12	-	45	72.6	339	1 O030640 METHYLCOBALAMIDE:COM MET	4.15e+01
13	45	72.6	339	1	Q48928 METHYLCOBALAMIDE:COM MET	4.15e+01
14	45	72.6	339	1	Q48950 METHYLCOBALAMIN: COENZ	4.15e+01
15	45	72.6	370	8	O48172 CYTOCHROME B.	4.15e+01
16	45	72.6	420	11	O08833 BILE ACID COA AMINO A	4.15e+01
17	45	72.6	420	11	O63276 KAN-L.	4.15e+01
18	45	72.6	980	5	Q175952 SIMILARITY TO INSULIN-	4.15e+01
19	45	72.6	1805	11	Q63667 FIBER 344 PRE-SIALOMU	4.15e+01
20	44	71.0	98	1	Q9728X HYPOTHETICAL 10.3 KD P	6.67e+01

			6.63e+01
		FLAGELLAR HOOK-ASSOCIA-	
		ABC TRANSPORTER, ATP-B	6.63e+01
		ACETYLCHOLINE RECEPTOR	1.05e+02
		ANTI-SIGMA F FACTOR AN	1.05e+02
		F13EF21.20 PROTEIN.	1.05e+02
		PHOSPHOGLYCOPROTEIN	1.05e+02
		PX01-66.	1.05e+02
		ARCA.	1.05e+02
		CYTOCROME P450 HYDRO-	1.05e+02
		HYPOTHETICAL 49.5 KD P	1.05e+02
		HYPOTHETICAL 53.2 KD P	1.05e+02
		W05H5.3 PROTEIN.	1.05e+02
		PUTATIVE ACYLTRANSFER-	1.05e+02
		PUTATIVE TRANSPEPTIDE.	1.05e+02
		RCH22 PROTEIN.	1.05e+02
		AUTOLYSIN SENSOR KINAS	1.05e+02
		HYPOTHETICAL 68.3 KD P	1.05e+02
		PUTATIVE SUGAR TRANSPO	1.05e+02
		PKS002C (FRAGMENT).	1.05e+02
		HYPOTHETICAL 165.6 KD	1.05e+02
		HYPOTHETICAL 166.6 KD	1.05e+02
		FK506 POLYPEPTIDE SYNTH	1.05e+02
		KDO SYNTHETASE.	1.65e+02
		Y102A5C.29 PROTEIN.	1.65e+02
		PARG PROTEIN.	1.65e+02
21	44	P94745	2
22	44	467	2
22	44	620	1
23	43	029198	
23	43	91	11
24	43	054712	
24	43	123	2
25	43	083551	
25	43	190	10
26	43	09XIA3	
26	43	243	2
26	43	086207	
27	43	2	
27	43	09X336	
28	43	2	
28	43	055855	
29	43	2	
29	43	09X5P8	
30	43	2	
30	43	068124	
31	43	2	
31	43	0509	2
32	43	2	
32	43	05457	2
33	43	2	
33	43	09X40	
33	43	579	5
33	43	09X7A9	
34	43	2	
34	43	053208	
35	43	2	
35	43	043390	
36	43	2	
36	43	049413	
37	43	2	
37	43	071749	
38	43	2	
38	43	065497	
38	43	729	10
39	43	2	
39	43	069497	
40	43	2	
40	43	050470	
41	43	3	
41	43	075727	
41	43	69.4	
41	43	075295	
42	43	2	
42	43	092685	
42	43	1616	2
42	43	092GA4	
43	42	2	
43	42	7576	2
43	42	092714	
43	42	269	2
43	42	67.7	
43	42	347	2
43	42	09XX84	
44	42	5	
44	42	046043	
45	42	768	5
45	42	67.7	

## ALIGNMENTS

RESULT	1	PRELIMINARY;	PRT;	766 AA.
ID	O23161;			
AC	O23161;	05' Created)		
DT	01-JAN-1998	(TREMBLrel. 05, Last sequence update)		
DT	01-MAY-1999	(TREMBLrel. 10, Last annotation update)		
DT	01-NOV-1999	(TREMBLrel. 12, Last annotation update)		
DE	RECEPTOR KINASE-LIKE PROTEIN (EC 2.7.1.1.)			
GN	C7A0..110.			
OS	Arabidopsis thaliana (Mouse-ear cress).			
OC	Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;			
OC	euphyllophytes; Spermato phyta; Magnoliophyta; eudicotyledons;			
OC	core eudicots; Rosidae; eurosids II; eurosids III; Brassicales; Brassicaceae;			
OC	Arabidopsis.			
RN	[1]			

```

Query Match          Score 53; DB 10; Length 766;
Best Local Similarity 70.0%; Pred. No. 7.44e-01;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0

335 DIAGIGILAY 344
:|||||:|
1 EARGIGILTY 10

RESULT          PRELIMINARY; PRT; 808 AA.
Q9ZT37;
Q9ZT37;
01-MAY-1999 (TREMBLE1. 10, Created)
01-MAY-1999 (TREMBLE1. 10, Last sequence update)
01-MAY-1999 (TREMBLE1. 10, Last annotation update)
PUTATIVE GLUTAMATE RECEPTOR.
GERL.

Arabidopsis thaliana (Mouse-ear cress).
Pukkarvara, Viridiniantha; Strennonhva: Tracheophyta;

```

OC euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;  
 OC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;  
 OC Arabidopsis.  
 [1]

RP SEQUENCE FROM N.A.  
 RX MEDLINE; 99039497.  
 RA LAM H.M., CHIU J., HSIEH M.H., MEISEL L., OLIVEIRA I.C., SHIN M.,  
 RA CORUZZI G.;  
 RT "Glutamate-receptor genes in plants.";  
 RL Nature 396:125-126(1998).  
 DR EMBL; AF079938; AAD09173.1; -.

KW Receptor.  
 SQ SEQUENCE 808 AA; 90518 MW; C3554B89 CRC32;

Query Match 3 PRELIMINARY; PRT; 165 AA.  
 ID P71810; LARDINOIS S., LAUBER J., LAZAREVIC V.,  
 AC P71810; LAPIDUS A., LARDINOIS S., LAUBER J., LAZAREVIC V.,  
 DT 01-FEB-1997 (TREMBLrel. 02, Created)  
 DT 01-NOV-1998 (TREMBLrel. 08, Last sequence update)  
 DE HYPOTHETICAL 18.2 KD PROTEIN.

GN MTCPB12.16.  
 OS Mycobacterium tuberculosis.

Bacteria; Firmicutes; Actinobacteria; Actinomycetales; Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.

[1] RP SEQUENCE FROM N.A.  
 RC STRAIN=H37RV;  
 RA BARREL B.G., RAJANDREAM M.A.;  
 RL Submitted (OCT-1996) to the EMBL/GenBank/DDJB databases.  
 RN [2]

RP SEQUENCE FROM N.A.  
 RC STRAIN=H37RV;  
 RA BARREL B.G., RAJANDREAM M.A.;  
 RL Submitted (OCT-1996) to the EMBL/GenBank/DDJB databases.  
 RN [3]

RP SEQUENCE FROM N.A.  
 RC STRAIN=H37RV;  
 RX MEDLINE; 98181548.

RA PHILIP W.J., POULET S., EIGELMEIER K., PASCOPELLA L.,  
 RA BALASUBRAMIAN V., HEYM B., BERGH S., BLOOM B.R., JACOBS W.R. JR.,  
 RA COLE S.T.;

RT "An integrated map of the genome of the tubercle bacillus,  
 RT Mycobacterium tuberculosis H37Rv, and comparison with Mycobacterium  
 RT leprae.";  
 RT Proc. Natl. Acad. Sci. U.S.A. 93:3132-3137(1996).

SQ SEQUENCE 165 AA; 18189 MW; BFB84C79 CRC32;

Query Match 4 PRELIMINARY; PRT; 250 AA.  
 Best Local Similarity 75.0%; Pred. No. 1.58e-01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

DR EMBL; Z81011; CAB02643.1; -.

KW Hypothetical protein.  
 SQ SEQUENCE 18189 MW; BFB84C79 CRC32;

Query Match 5 PRELIMINARY; PRT; 478 AA.  
 Best Local Similarity 75.0%; Pred. No. 1.58e-01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

DR EMBL; Z99110; CAB12998.1; -.

RC SEQUENCE 250 AA; 30119 MW; C96222FD CRC32;

Query Match 6 PRELIMINARY; PRT; 1.58e+01 AA.  
 Best Local Similarity 66.7%; Pred. No. 1.58e+01;  
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

DR 89 TDGIGILAV 97  
 : |||||:|  
 Qy 2 AAGIGILIV 10

RESULT 5  
 ID Q87090  
 AC Q87090;  
 DT 01-NOV-1996 (TREMBLrel. 01, Created)

DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)

OS Glycoprotein GII.

Pseudorabies virus.

OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;

OC Alphaherpesvirinae; Varicellovirus.

RN [1]  
 RP SEQUENCE FROM N.A.

RC STRAIN=INDIANA\_S;

RX MEDLINE; 961316347.

RA ISHIKAWA K., TSURUI M., TAGUCHI K., SAITO H., MURAMATSU M.;

DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)  
 DE YJBA PROTEIN.

GN YJBA.

OS Bacillus subtilis.

OC Bacteria; Firmicutes; Bacillus/Clostridium group;

OC Bacillus/Staphylococcus group; Bacillus.

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=168;

RC MEDLINE; 98044033.

RA KUNST F., OGASAWARA N., MOSZER L., ALBERTINI A.M., ALLONI G.,

RA AZEVEDO V., BERPERO M.G., BESSIERES P., BOLOTIN A., BORCHERT S.,

RA BORRISS R., BOURSET L., BRANS A., BRAUN M., BRIGNELL S.C., BRON S.,

RA BROUILLET S., BRUSCHI C.V., CALDWELL B., CAPUANO V., CARTER N.M.,

RA CHOI S.K., CODANI J.J., CONNERTON T.F., CUMMINGS N.J., DANIEL R.A.,

RA DENIZOT F., DEVINE K.M., DUSTERRHOFT A., EHRLICH S.D., ENMRISON P.T.,

RA ENTIAN K.D., ERINGTON J., FABRET C., FERRARI E., FOULGER D.,

RA FRITZ C., FUJITA M., FUJITA Y., FUMA S., GALIZZI A., GALLERON N.,

RA GUISEPEI G., GUY B.J., GOFFEAU A., GOLIGHTY E.J., GRANDI G.,

RA HILBERT H., HOLSSAPPEL S., HOSONO S., HULLO M.F., ITAYA M., JONES L.,

RA JORIS B., KARAMATA D., KASHARA Y., KLAERF-BLANCHARD M., KLEIN C.,

RA KOBAYASHI Y., KOETTER P., KONINGSTEIN G., KROGH S., KUMANNO M.,

RA KURITA K., LAPIDUS A., LARDINOIS S., LAUBER J., LAZAREVIC V.,

RA LEE S.M., LEVINE A., LIU H., MASUDA S., MAUEL C., MEDIGUE C.,

RA MEDINA N., MELLADO R.P., MIZOUMA M., MOESTL D., NAKAI S., NOBACK M.,

RA NOONE D., O'REILLY M., OGAWA K., OGAWA K., OGAWA K., OGAWA K.,

RA PARRO V., POHL T.M., PORWOLLIK S., PRESCOTT A.M.,

RA PRESECAN E., PUJIC P., PURNELLE D., RAPORT G., REY M., REYNOLDS S.,

RA RIEGER M., RIVOLTA C., ROCHA E., ROCHE B., ROSE M., SADALE E.Y.,

RA SATO T., SCANLAN E., SCHLEICH S., SCROFONE F.,

RA SEKIGUCHI J., SEKOWSKA A., SEROR S.J., SEROR P.P., SHIN B.S., SOLDO B.,

RA SOROKIN A., TACCONI E., TAKAGI T., TAKAHASHI H., TAKEMARU K.,

RA TAKEUCHI M., TAMAKOSHI A., TANAKA T., TERESTRA P., TOGNONI A.,

RA TOSATO V., UCHIYAMA S., VANDENBOL M., VANNIER F., VASSAROTTI A.,

RA VIARI A., WAMBUT R., WEDLER E., WEDLER H., WEITZENEGGER T.,

RA WINTERS P., WIPAT A., YAMAMOTO H., YAMANE K., YATA K.,

RA YOSHIDA K., YOSHIKAWA H.F., ZUMSTEIN E., YOSHIKAWA H.,

RA "The complete genome sequence of the gram-positive bacterium Bacillus

RT subtilis.";

RL Nature 390:249-256(1997).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=168;

RA KUNST F., OGASAWARA N., YOSHIKAWA H., DANCHIN A.,

RA TOSATO V., UCHIYAMA S., VANDENBOL M., VANNIER F., VASSAROTTI A.,

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RA VIARI A., WAMBUT R., WEDLER E., WEDLER H., WEITZENEGGER T.,

RA WINTERS P., WIPAT A., YAMAMOTO H., YAMANE K., YATA K.,

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RN [2]

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RA KUNST F., OGASAWARA N., YOSHIKAWA H., DANCHIN A.,

RA TOSATO V., UCHIYAMA S., VANDENBOL M., VANNIER F., VASSAROTTI A.,

RA VIARI A., WAMBUT R., WEDLER E., WEDLER H., WEITZENEGGER T.,

RA WINTERS P., WIPAT A., YAMAMOTO H., YAMANE K., YATA K.,

RA YOSHIDA K., YOSHIKAWA H.F., ZUMSTEIN E., YOSHIKAWA H.,

RA "The complete genome sequence of the gram-positive bacterium Bacillus

RT subtilis.";

RL Nature 390:249-256(1997).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=168;

RA KUNST F., OGASAWARA N., YOSHIKAWA H., DANCHIN A.,

RA TOSATO V., UCHIYAMA S., VANDENBOL M., VANNIER F., VASSAROTTI A.,

RA VIARI A., WAMBUT R., WEDLER E., WEDLER H., WEITZENEGGER T.,

RA WINTERS P., WIPAT A., YAMAMOTO H., YAMANE K., YATA K.,

RA YOSHIDA K., YOSHIKAWA H.F., ZUMSTEIN E., YOSHIKAWA H.,

RA "The complete genome sequence of the gram-positive bacterium Bacillus

RT subtilis.";

RL Nature 390:249-256(1997).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=168;

RA KUNST F., OGASAWARA N., YOSHIKAWA H., DANCHIN A.,

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RA VIARI A., WAMBUT R., WEDLER E., WEDLER H., WEITZENEGGER T.,

RA WINTERS P., WIPAT A., YAMAMOTO H., YAMANE K., YATA K.,

RA YOSHIDA K., YOSHIKAWA H.F., ZUMSTEIN E., YOSHIKAWA H.,

RA "The complete genome sequence of the gram-positive bacterium Bacillus

RT subtilis.";

RL Nature 390:249-256(1997).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=168;

RA KUNST F., OGASAWARA N., YOSHIKAWA H., DANCHIN A.,

RA TOSATO V., UCHIYAMA S., VANDENBOL M., VANNIER F., VASSAROTTI A.,

RA VIARI A., WAMBUT R., WEDLER E., WEDLER H., WEITZENEGGER T.,

RA WINTERS P., WIPAT A., YAMAMOTO H., YAMANE K., YATA K.,

RA YOSHIDA K., YOSHIKAWA H.F., ZUMSTEIN E., YOSHIKAWA H.,

RA "The complete genome sequence of the gram-positive bacterium Bacillus

RT subtilis.";

RL Nature 390:249-256(1997).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=168;

RA KUNST F., OGASAWARA N., YOSHIKAWA H., DANCHIN A.,

RA TOSATO V., UCHIYAMA S., VANDENBOL M., VANNIER F., VASSAROTTI A.,

RA VIARI A., WAMBUT R., WEDLER E., WEDLER H., WEITZENEGGER T.,

RA WINTERS P., WIPAT A., YAMAMOTO H., YAMANE K., YATA K.,

RA YOSHIDA K., YOSHIKAWA H.F., ZUMSTEIN E., YOSHIKAWA H.,

RA "The complete genome sequence of the gram-positive bacterium Bacillus

RT subtilis.";

RL Nature 390:249-256(1997).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=168;

RA KUNST F., OGASAWARA N., YOSHIKAWA H., DANCHIN A.,

RT "Sequence variation of the gc gene among pseudorabies virus strains.";  
 RL VET. MICROBIOL 49:267-272(1996).  
 DR EMBL: D49436; BAA08414.1.  
 DR PRINTS; PRO0668; GLYCPROTEIN.

SEQUENCE 478 AA; 51150 MW; D6A143B4 CRC32;

Query Match Score 47; DB 14; Length 478;  
 Best Local Similarity 75.08; Pred. No. 1.58e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 DR 455 AGIGILAI 462  
 |||||||:  
 QY 3 AGIGILTV 10

RESULT 6 PRELIMINARY; PRT; 479 AA.  
 ID Q87089; PRELIMINARY; PRT; 479 AA.  
 AC Q87089; PRELIMINARY; PRT; 479 AA.  
 DT 01-NOV-1996 (TREMBLrel. 01, Created)  
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)  
 DE GLYCOPROTEIN GTII.  
 OS Pseudorabies virus.  
 VS dsDNA viruses, no RNA stage; Herpesviridae;  
 CC Alphaherpesvirinae; varicellovirus.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=YAMAGATA S-81;  
 RM MEDLINE; 96316347.

RA ISHIKAWA K.; TSUTSUI M.; TAGUCHI K.; SAITOH A.; MURAMATSU M.;  
 RT "Sequence variation of the gc gene among pseudorabies virus strains.";  
 RL VET. MICROBIOL. 49:267-272(1996).  
 DR EMBL: D49435; BAA08413.1.  
 DR PRINTS; PRO0668; GLYCPROTEIN.

SEQUENCE 479 AA; 51109 MW; A009EB9B CRC32;

Query Match Score 47; DB 14; Length 479;  
 Best Local Similarity 75.08; Pred. No. 1.58e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 DR 456 AGIGILAI 463  
 |||||||:  
 QY 3 AGIGILTV 10

RESULT 7 PRELIMINARY; PRT; 479 AA.  
 ID Q87091; PRELIMINARY; PRT; 479 AA.  
 AC Q87091; PRELIMINARY; PRT; 479 AA.  
 DT 01-NOV-1996 (TREMBLrel. 01, Created)  
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)  
 DE GLYCOPROTEIN GTII.  
 OS Pseudorabies virus.  
 VS dsDNA viruses, no RNA stage; Herpesviridae;  
 CC Alphaherpesvirinae; varicellovirus.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=NIA3;

RM MEDLINE; 96316347.  
 RA ISHIKAWA K.; TSUTSUI M.; TAGUCHI K.; SAITOH A.; MURAMATSU M.;  
 RT "Sequence variation of the gc gene among pseudorabies virus strains.";  
 RL VET. MICROBIOL. 49:267-272(1996).  
 DR EMBL: D49437; BAA08415.1.  
 DR PRINTS; PRO0668; GLYCPROTEIN.

SEQUENCE 479 AA; 51148 MW; CC3EEFF9A CRC32;

? Query Match Score 47; DB 14; Length 479;  
 Best Local Similarity 75.08; Pred. No. 1.58e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 DR 456 AGIGILAI 463  
 |||||||:  
 QY 3 AGIGILTV 10

RESULT 8 PRELIMINARY; PRT; 773 AA.  
 ID O22178; PRELIMINARY; PRT; 773 AA.  
 AC O22178; PRELIMINARY; PRT; 773 AA.  
 DT 01-JAN-1998 (TREMBLrel. 05, Created)  
 DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)  
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)  
 DE PUTATIVE RECEPTOR PROTEIN KINASE.  
 GN T20016.7.  
 OS Arabidopsis thaliana (Mouse-ear cress).  
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;  
 OC euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;  
 OC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;  
 OC Arabidopsis.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=CV. COLUMBIA;  
 RA ROUNSLAY S.-D.; LIN X.; KETCHUM K.A.; CROSBY M.L.; BRANDON R.C.;  
 RA SYKES S.M.; KAUL S.; MASON T.M.; KERLAYAGE A.R.; ADAMS M.D.;  
 RA SOMERVILLE C.R.; VENTER J.C.;  
 RL Submitted (NOV-1997) to the EMBL/GenBank/DBJ databases.  
 EMBL: AC002391; AAB87101.1;  
 DR MENDEL; PF00506; Arath; 3435; 25106.  
 DR PFAM; PF00560; LRR; 4.  
 DR PFAM; PF00069; pkinase\_1.  
 DR SEQ SEQUENCE 773 AA; 8418 MW; 83C3953B CRC32;  
 DR Query Match Score 47; DB 10; Length 773;  
 DR Best Local Similarity 60.0%; Pred. No. 1.58e+01;  
 DR Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;  
 DR 341 DIAGITLAL 350  
 :||||||:  
 QY 1 EAAGGILTY 10

RESULT 9 PRELIMINARY; PRT; 2219 AA.  
 ID Q23388; PRELIMINARY; PRT; 2219 AA.  
 AC Q23388; PRELIMINARY; PRT; 2219 AA.  
 DT 01-NOV-1996 (TREMBLrel. 01, Created)  
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
 DT 01-JAN-1999 (TREMBLrel. 09, Last annotation update)  
 DE ZK1067.2 PROTEIN.  
 GN ZK1067.2.  
 OS Caenorhabditis elegans.  
 OC Eukaryota; Metazoa; Nemataoda; Secernentea; Rhabditia; Rhabditida;  
 OC Rhabditina; Rhabditoidea; Rhabditidae; Peledorinae; Caenorhabditis.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RA THOMAS K.;  
 RL Submitted (MAR-1996) to the EMBL/GenBank/DBJ databases.  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE: 94150718.  
 RA WILSON R.; AINSCOUGH R.; ANDERSON K.; BAYNES C.; BERKS M.,  
 RA BONFIELD J.; BURTON J.; CONNELL M.; COOPER J.; COULSON A.,  
 RA CRAYTON M.; DEAR S.; DU Z.; DURBIN R.; FAVELLO A.; FULTON L.,  
 RA GARDNER A.; GREEN P.; HAWKINS T.; HILLIER L.; JIER M.; JOHNSTON L.,  
 RA JONES M.; KERSHAW J.; KIRSTEN J.; LAISTER N.; LATREILLE P.,  
 RA LIGHTNING J.; LLOYD C.; MC MURRAY A.; MORTIMORE B.; O'CALLAGHAN M.,  
 RA PARSONS J.; PERCY C.; RIFKEN L.; ROOPRA A.; SAUNDERS D.; SHOWKEN R.,  
 RA SMAULDON N.; SMITH A.; SONNHAMMER E.; STADEN R.; SULSTON J.,  
 RA THERRY-MIEG J.; THOMAS K.; VAUDIN M.; VAUGHAN K.; WATERSTON R.,  
 RA WATSON A.; WEINSTOCK L.; WILKINSON-SPROAT J.; WOOLDRMAN P.;  
 RT RT elegans";  
 RL Nature 368:32-38(1994).  
 DR DR SEQUENCE Z7003B; CAA93884.1;  
 SQ SEQUENCE 2219 AA; 253649 MW; 59DE8B43 CRC32;  
 DR Query Match Score 47; DB 5; Length 2219;  
 DR Best Local Similarity 60.0%; Pred. No. 1.58e+01;



Query Match 72.6%; Score 45; DB 1; Length 339;  
 Best Local Similarity 75.0%; Pred. No. 4.15e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 Db 288 GIGILAV 294  
 |||||:  
 Qy 4 GIGILAV 10

Search completed: Fri May 5 22:11:45 2000  
 Job time : 89 secs.

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RESULT 14 PRELIMINARY; PRT; 339 AA.

ID Q48950; AC Q48950; DT 01-NOV-1996 (TREMBLrel. 01, Created)  
 RX MEDLINE; 96184544.  
 RA HARMS U.; THAUER R.K.;  
 RT "Methylcobalamin: coenzyme M methyltransferase isoenzymes MtaA and  
 MtbA from Methanosarcina barkeri. Cloning, sequencing and differential  
 transcription of the encoding genes, and functional overexpression of  
 the mtaA gene in Escherichia coli.";  
 RL Eur. J. Biochem. 235:653-659(1996).  
 DR EMBL; X91894; CA62996; 1.  
 DR PFAM; PF01208; URO-D; 1.  
 RW Transferase; Methyltransferase; DRUG-D; 1.  
 SEQUENCE 339 AA; 36761 MW; 5F6F0A9C CRC32;

Query Match 72.6%; Score 45; DB 1; Length 339;  
 Best Local Similarity 75.0%; Pred. No. 4.15e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 Db 307 AGYGLITV 314  
 |||||:  
 Qy 3 AGIGILTV 10

---

RESULT 15 PRELIMINARY; PRT; 370 AA.

ID Q48172; AC Q48172; DT 01-JUN-1998 (TREMBLrel. 06, Created)  
 RX MEDLINE; 98120004.  
 RA ANTARAMIAN A.; FUNES-ARGUELLO S.; VAZQUEZ-ACEVEDO M.; CORIA R.,  
 RA GONZALES-HALPHEN D.;  
 HL Submitted (JAN-1997) to the EMBL/GenBank/DDBJ databases.  
 COB.  
 OS Polytomella sp. 'Pringsheim 198-80'.  
 OG Mitochondrion.  
 OC Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; volvocales;  
 OC Chlamydomonadaceae; Polytomella.  
 RN SEQUENCE FROM N.A.  
 RC STRAIN=198-80, FROM E.G. PRINGSHEIM;  
 RA ANTARAMIAN A.; FUNES-ARGUELLO S.; VAZQUEZ-ACEVEDO M., CORIA R.,  
 RA GONZALES-HALPHEN D.;  
 HL Submitted (JAN-1997) to the EMBL/GenBank/DDBJ databases.  
 DR EMBL; U87396; AC24896; 1.  
 DR MENDER; 23585; P0538; P05385.  
 DR PFAM; PF00032; cytochrome\_b\_c\_1.  
 DR PFAM; PF00033; cytochrome\_b\_N\_1.  
 RW Mitochondrion.  
 SEQUENCE 370 AA; 41226 MW; 5D617081 CRC32;

Query Match 72.6%; Score 45; DB 8; Length 370;  
 Best Local Similarity 85.7%; Pred. No. 4.15e+01;  
 Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

(TM) 

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protein - protein database search, using Smith-Waterman algorithm  
psrclpp run on: Fri May 5 22:09:18 2000; MasPar time 3.01 Seconds

99.300 Million cell updates/sec  
abular output not generated.

title: >US-09-267-439-17  
description: (1-10) from US09267439.pep

perfect score: 62  
sequence: 1 EAAGIGILTV 10

scoring table:

searched: 82229 seqs, 29864866 residues

post-processing: Minimum Match 0% Listing first 45 summaries

database: swiss-prot38  
-

statistics: Mean 23.268; Variance 24.186; scale 0.962  
I:swissprot

Pred. No. is the number of results predicted by chance to have a

and is derived by analysis of the total score distribution.

8 Query result

No.	Score	Match	Length	DB ID	Description	Pred.	No.
1	6.2	100.0	118	1	MARI HUMAN MELANOMA ANTIGEN RECOG	8	83e-04

2	4.8	77.4	332	1
3	4.7	75.8	101	1
4	4.7	75.0	110	1
				ATPL-SULAC MEMBRANE-ASSOCIATED AT 5.22E+00 5.22E+00 5.22E+00 5.22E+00

4	4.7	75.8	110	VQAD_LAMBDD
4	4.7	75.8	384	PQQ_METEX
6	4.7	75.8	479	VGLC_PRVIF
				HEAD DECORATION PROTE
				COENZYME PQQ SYNTHESIS
				GLYCOPROTEIN GII PREC
				5.22E+00
				5.22E+00

9	46	4/4	2/1	YK23_YEAST	HYPOTHETICAL	31.0	KD P	8.7e+00
10	45	72.6	201	1 Y760_RHISN	HYPOTHETICAL PROTEIN	P	1.46e+01	
11	45	72.6	291	1 Y4TQ_RHISN	PROBABLE PEPTIDE ABC T	A	1.46e+01	

4.4	71.0	409	PLID-BCUL1	ENDO-1,4-BETA-XYLANASE	2.41e+01
4.4	71.0	635	XYND-PAEPO	ENDO-1,4-BETA-XYLANASE	2.41e+01
4.3	69.4	132	ATPE_ARATH	ATP SYNTHASE EPSILON C	3.94e+01
4.3	69.4	132	ATPE_ARATH	ATP SYNTHASE EPSILON C	3.94e+01

•1.7	43	69.4	207	1 RAP-ECOLI 50S RIBOSOMAL PROTEIN
1.8	43	69.4	345	1 TAP-ECOLI ALKALINE PHOSPHATASE I
1.9	43	69.4	404	1 SGAA-HPMW SERINE-GLYOXYLATE AMYLASE
2.0	43	69.4	61	1 C12H22O11

4.0	4.3	69.4	493	1	YACC_BACSU	HYPOTHALAMIC METABOLIT	3.94e+01
2.1	4.3	69.4	493	1	ACHE_MOUSE	ACETYLCHOLINE RECEPTOR	3.94e+01
2.2	4.3	69.4	590	1	CHL1_ARATH	NITRATE/CHLORATE TRANS	3.94e+01

24	651	1	BGLR_CANFA	BETA-GLUCURONIDASE PRE-	3.94e+01
43	675	1	NUML_ACACA	NADH DEHYDROGENASE F	3.94e+01
43	1217	1	EGR_MOUSE	PRO-EPIDEMYL GROWTH F	3.94e+01
43	1325	1	YDEK_ECOLI	HYPOTHETICAL 136.5 KD	3.94e+01
43	111	1	YPEE_ECOLI	HYPOTHETICAL 11.6 KD P	6.37e+01
42	67	7	FLA2_METHV	FLAGELLIN B2 PRECURSOR	6.37e+01
42	67	7	FLA2_METHV	FLAGELLIN B2 PRECURSOR	6.37e+01
42	67	22	FLA2_METHV	FLAGELLIN B2 PRECURSOR	6.37e+01
42	251	1	YFQF_ECOLI	HYPOTHETICAL TRANSCRIPT	6.37e+01
42	308	1	YFQA_HARIN	1,4-DIHYDROXY-2-NAPHTH	6.37e+01
42	321	1	MENA_HARIN	HYPOTHETICAL 35.0 KD P	6.37e+01
42	325	1	RCEM_CHRV1	REACTION CENTER PROTEIN	6.37e+01
42	67	34	HRCA_SPMU	HEAT INDUCIBLE TRANSFER	6.37e+01
42	67	35	HRCA_SPMU	PUTATIVE UDP-GLICURONO	6.37e+01
42	67	44	UGTC_CABEL	PUTATIVE THIOPHENYL AND	6.37e+01
42	67	44	UGTC_CABEL	LYSINE TRANSPORT PRO	6.37e+01
42	67	44	UGTC_CABEL	PROTEIN-EXPORT MEMBRAN	6.37e+01
42	67	36	THDF_HAEIN	ACIN INTERACTING PROT	6.37e+01
42	67	37	THDF_HAEIN	HYPOTHETICAL 74.3 KD P	6.37e+01
42	67	501	LYSI_CORG1	NADH-DEPENDENT FLAVIN	6.37e+01
42	67	503	SEED_HELPJ	PUTATIVE MEMBRANE PROT	6.37e+01
42	67	526	SEED_HELPJ	BREFELDIN A RESISTANCE	6.37e+01
42	67	530	AIP2_YEAST	BREFELDIN B1	6.37e+01
42	67	530	AIP2_YEAST	SCHIPO	6.37e+01
42	67	659	YB7T_BACSU	YB7T_BACSU	6.37e+01
42	661	1	BATH_EUBSP	YODH_BACSU	6.37e+01
42	67	7	YODH_BACSU	YODH_BACSU	6.37e+01
42	1530	1	BFRP1_SCIPHO	BFRP1_SCIPHO	6.37e+01

ALIGNMENTS					
RESULT	1	MARI_HUMAN	STANDARD;	PRT;	118 AA.
ID		Q16655;			
AC		Q16655;			
DT	01-NOV-1997	(Rel. 35, Created)			
DT	01-NOV-1997	(Rel. 35, Last sequence update)			
DT	15-JUL-1998	(Rel. 36, Last annotation update)			
DE	(ANTIGEN SK29-AA) (ANTIGEN LB39-AA).				
GN	MELANA OR MART1.				
OS	Homo sapiens (Human).				
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;				
OC	Eutheria; Primates; Catarrhini; Hominidae; Homo.				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RC	TISSUE=MESENTERIC MELANOMA;				
RX	MEDLINE: 94224770.				
RA	KAWAKAMI Y., ELIYAHU S., DELGADO C.H., ROBBINS P.F., RIVOLTINI L.,				
RA	TOPALIAN S.L., MIKI T., ROSENBERG S.A.;				
RT	"Cloning of the gene coding for a shared human melanoma antigen recognized by autologous T cells infiltrating into tumor.";				
RT	Proc. Natl. Acad. Sci. U.S.A. 91:515-519(1994).				
RL	[2]				
RN					
RP	SEQUENCE FROM N.A.				
RX	MEDLINE: 94273389.				
RA	COULIE P.G., BRICHARD V., VAN PEL A., WOELFEL T., SCHNEIDER J.J.,				
RA	TRAVERSARI C., BOON T., DE PLAEN E., LURQUIN C., SZIKORA J.-P.,				
RA	RENAUD J.-C., BOON T.;				
RT	"A new gene coding for a differentiation antigen recognized by autologous cytolytic T lymphocytes on HLA-A2 melanomas.";				
RT	J. Exp. Med. 180:35-42(1994)				
RL	-1 - TISSUE SPECIFICITY: EXPRESSION IS RESTRICTED TO MELANOMA AND MELANOCTY CELL LINES AND RETINA.				
CC					
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CC	-				
CC	EMBL; U06452; AAA19238; 1; -.				
CC	EMBL; U06654; AAA20389; 1; -.				
CC	Antigen; Transmembrane.				
KW					
FT	POTENTIAL.				
SO	27				
SO	118 AA; 13157 MW; DFF22CFCS CRC32;				

ALIGNMENTS

RESULT	1	MARI HUMAN	STANDARD;	PRT;	118 AA.
ID	Q16655				
AC	Q16655;1997	(Rel. 35, Created)			
DT	01-NOV-1997	(Rel. 35, Last sequence update)			
DT	01-NOV-1997	(Rel. 36, Last annotation update)			
DT	15-JUL-1998	(Rel. 36, Last annotation update)			
DE	MELANOMA ANTIGEN RECOGNIZED BY T-CELLS 1 (MART-1) (MELAN-A PROTEIN)				
DE	(ANTIGEN SK29-AA) (ANTIGEN LB39-AA).				
GN	MELANA OR MART1.				
OS	Homo sapiens (Human).				
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;				
OC	Eutheria; Primates; Catarrhini; Hominidae; Homo.				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RC	TISSUE=MELANOMA;				
RX	MEDLINE; 94224770.				
RA	KAWAKAMI Y., ELIYAHU S., DELGADO C.H., ROBBINS P.F., RIVOLTINI L.,				
RA	TOPALIAN S.L., MIKI T., ROSENBERG S.A.;				
RT	Cloning of the gene coding for a shared human melanoma antigen recognized by autologous T cells infiltrating into tumor.;				
RT	Proc Natl Acad Sci U S A 1994 Jan 25;91:2515-2519.				

[RN 1]  
RN SEQUENCE FROM N.A.  
RP MEDLINE; 94:275389.  
RX  
RA COUGL P.G., BRICHARD V., VAN PEL A., WOELFEL T., SCHNEIDER J.,  
RA TRAVERSARI C., MATEI S., DE PLAEN E., LURQUIN C., SZIKORA J.-P.,  
RA RENAULD J.-C., BOON T.;  
RT "A new gene coding for a differentiation antigen recognized by  
RT autologous cytolytic T lymphocytes on HLA-A2 melanomas.";  
RL J. Exp. Med. 180:35-42 (1994).  
CC - - TISSUE SPECIFICITY: EXPRESSION IS RESTRICTED TO MELANOMA AND  
CC MELANOCYTE CELL LINES AND RETINA.  
CC  
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CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).  
CC  
CC EMBL: U06452; AAA19238.1; -;  
DR EMBL: U06654; AAA20389.1; -;  
KW Antigen; Transmembrane.  
FT 27 47 POTENTIAL.  
SO SEQUENCE  
DR 118 AA: 113157 MW: DFF2CF6 CRC32;  
SO 118 AA: 113157 MW: DFF2CF6 CRC32;

Query Match 100.0%; Score 62; DB 1; Length 118;  
 Best Local Similarity 100.0%; Pred. No. 8.83e-04;  
 NMatches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Sulfolobus acidocaldarius. Crenarchaeota; Sulfolobales; Sulfolobus.  
 [1]  
 OS OC RN  
 Sequence from N.A., AND PARTIAL SEQUENCE.  
 RP RX  
 MEDLINE: 89214142.  
 RA DENDA K., KONISHI J., OSHIMA T., DATE T., YOSHIDA M.;  
 RT "A gene encoding the proteolipid subunit of *Sulfolobus acidocaldarius*  
 ATPase complex."  
 RT RT  
 RL J. Biol. Chem. 264:7119-7121(1989).

RESULT 2 ID ACOA\_ALCEU STANDARD; PRT; 332 AA.  
 AC P27745;  
 DT 01-AUG-1992 (Rel. 23, Created)  
 DT 01-DEC-1992 (Rel. 24, Last sequence update)  
 DT 01-DEC-1992 (Rel. 24, Last annotation update)  
 DE ACETOIN:2,6-DICHLOROPHENOL OXIDOREDUCTASE ALPHA  
 SUBUNIT (EC 1.1.1.-) (ACETOIN:DCPIP OXIDOREDUCTASE-ALPHA)  
 DE (AO:DCPIP OR).  
 GN ACOA.  
 Alcaligenes eutrophus. Proteobacteria; beta subdivision; Burkholderia group;  
 Ralstonia.  
 OC RN  
 Sequence from N.A., AND SEQUENCE OF 1-31.  
 RC STRAIN=H16;  
 RX MEDLINE: 91286190.  
 RA PRIEBERT H., HEIN S., KRUGGER N., ZEH K., SCHMIDT B., STEINBUCHEL A.;  
 RT "Identification and molecular characterization of the Alcaligenes  
 eutrophus H16 aco operon genes involved in acetoin catabolism.";  
 RL J. Bacteriol. 173:4056-4071(1991).  
 CC -1- FUNCTION: CATALYZES THE 2,6-DICHLOROPHENOL-DEPENDENT  
 CLEAVAGE OF ACETOIN INTO ACETATE AND ACETALDEHYDE. IN VITRO, THE  
 ALPHA SUBUNIT IS PROBABLY THE CATALYTIC SUBUNIT OF THE ENZYME.  
 CC -1- COFACTOR: THIAMINE PYRROPHOSPHATE.  
 CC -1- PATHWAY: ACETOIN CATABOLISM.  
 CC -1- SUBUNIT: TETRAMER OF TWO ALPHA AND TWO BETA SUBUNITS.  
 CC -1- SIMILARITY: TO THE ALPHA SUBUNITS OF 2-OXO-ACID DEHYDROGENASE  
 COMPONENTS OF VARIOUS MULTIZYME COMPLEXES.  
 CC -1-  
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 or send an email to license@isb-sib.ch).  
 CC -1-  
 DR EMBL; M66060; AAA21948.1; -.  
 DR PIR; B42462; DEALXKE.  
 DR PFAM; PF00676; E1\_dehydrog; 1.  
 KW Oxidoreductase; Flavoprotein; thiamine pyrophosphate.  
 FT BINDING 0 0  
 SQ SEQUENCE 332 AA; 35243 MW; 332DA8E CRC32; THIAMINE PYROPHOSPHATE (BY SIMILARITY).

Query Match 77.4%; Score 48; DB 1; Length 332;  
 Best Local Similarity 87.5%; Pred. No. 3.08e+00;  
 NMatches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Sulfolobus acidocaldarius. Crenarchaeota; Sulfolobales; Sulfolobus.  
 [1]  
 OS OC RN  
 Sequence from N.A., AND PARTIAL SEQUENCE.  
 RP RX  
 MEDLINE: 89214142.  
 RA DENDA K., KONISHI J., OSHIMA T., DATE T., YOSHIDA M.;  
 RT "A gene encoding the proteolipid subunit of *Sulfolobus acidocaldarius*  
 ATPase complex."  
 RT RT  
 RL J. Biol. Chem. 264:7119-7121(1989).

RESULT 3 ID ATPL\_SULAC STANDARD; PRT; 101 AA.  
 AC P23040;  
 DT 01-NOV-1991 (Rel. 20, Created)  
 DT 01-NOV-1991 (Rel. 20, Last sequence update)  
 DE MEMBRANE ASSOCIATED ATPASE C CHAIN (EC 3.6.1.34) (SUL-ATPASE  
 DE PROTEOLIPID CHAIN).  
 GN ATPP.

CC -1- FUNCTION: SPABILIZES THE HEAD SHELL FOLLOWING THE REARRANGEMENT  
 OF THE GPE SUBUNITS OF THE HEAD SHELL LATTICE THAT ACCOMPANIES  
 EXPANSION OF THE HEAD. THERE ARE APPROXIMATELY 420 COPIES OF  
 PROTEIN D PER MATURE PHAGE.  
 CC -1- SIMILARITY: TO BACTERIOPHAGE 21 HEAD DECORATION PROTEIN.

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DR	EMBL; J02459; AAA6539.1;	-.
DR	PIR; A04334; VHBPD1.	.
DR	KW	Coat protein.
DR	PIR; A3206; A23206.	.
DR	SEQUENCE 110 AA;	11572 MW; FDD50011 CRC32;
DR	Query Match 5	Score 47; DB 1; Length 110;
	Best Local Similarity 50.0%;	Pred. No. 5.22e+00;
DR	Matches 5;	Mismatches 0; Indels 0; Gaps 0;
DR	Conservative 5;	
DR	Db 55 DGAVGILAV 64	
DR	::: : : : :	
DR	Dy 1 EAAGIGILTV 10	
RESULT	5 PQQE_METEX	STANDARD; PRT; 384 AA.
ID	AC P71517;	
RA	TOYAMA H., CHISTOSERDOVA L., LIDSTRÖM M.E.;	
RT	Sequence analysis of pqqe genes required for biosynthesis of pyrrolequinoline quinone in Methylobacterium extorquens AM1 and the purification of a biosynthetic intermediate.;	
RT	Microbiology 143:595-602(1997).	
RL	-1- FUNCTION: REQUIRED FOR COENZYME PYRROLO-QUINOLINE-QUINONE (PQQ) BIOSYNTHESIS.	
SN	-1- SIMILARITY: BELONGS TO THE MOAA/NIFB/PQOE FAMILY OF PROTEINS.	
DS	Methylobacterium extorquens.	
OC	Bacteria; Proteobacteria; alpha subdivision; Methylobacterium.	
RN	[1]	
RP	SEQUENCE FROM N.A.	
RC	STRAINNAME / NCIB 9133;	
RX	MEDLINE; 97195805.	
RA	TOYAMA H., CHISTOSERDOVA L., LIDSTRÖM M.E.;	
RT	Sequence analysis of pqqe genes required for biosynthesis of pyrrolequinoline quinone in Methylobacterium extorquens AM1 and the purification of a biosynthetic intermediate.;	
RT	Microbiology 143:595-602(1997).	
RL	-1- FUNCTION: REQUIRED FOR COENZYME PYRROLO-QUINOLINE-QUINONE (PQQ) BIOSYNTHESIS.	
CC	-1- SIMILARITY: BELONGS TO THE MOAA/NIFB/PQOE FAMILY OF PROTEINS.	
IC	EMBL; U72662; AAB58898.1;	.
IC	PROSTIE; PS01305; MOAA_NIFB_PQQE; 1.	.
IC	PFAAM; PF01444; Moaa_NifB_Pqqe; 1.	.
IC	PQOE; Iron-sulfur.	.
IC	METAL 28 28 IRON-SULFUR (POTENTIAL).	.
IC	METAL 32 32 IRON-SULFUR (POTENTIAL).	.
IC	METAL 35 35 IRON-SULFUR (POTENTIAL).	.
IC	SEQUENCE 384 AA; 41714 MW; 7BD3BE8C CRC32;	.
IC	Query Match 5	Score 47; DB 1; Length 384;
	Best Local Similarity 70.0%;	Pred. No. 5.22e+00;
IC	Matches 7;	Mismatches 1; Indels 0; Gaps 0;
IC	Conservative 7;	
IC	Dy 59 EAAGIGLVHV 68	
IC	: : :	
IC	Dy 1 EAAGIGILTV 10	

RESULT	6	VGLPFV_PRT	STANDARD;	PRT;	479 AA.
ID	PO6024;				
AC	13-AUG-1987 (Rel. 05, Created)				
DT	01-APR-1993 (Rel. 25, Last annotation update)				
DT	GLYCOPROTEIN GII.1 PRECURSOR.				
OS	Pseudorabies virus (strain Indiana-Funkhauser / Becker) (PRV).				
OC	Viruses; dsDNA viruses, no RNA stage; Herpesviridae;				
OC	Alphaherpesvirinae; Varicellovirus.				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RX	MEDLINE: BG200375.				
RA	ROBBINS A.K., WATSON R.J., WHEALY M.E., HAYS W.W., ENQUIST L.W.;				
RT	"Characterization of a pseudorabies virus glycoprotein gene with homology to herpes simplex virus type 1 and type 2 glycoprotein C."				
RL	J. Virol. 58:339-347(1986).				
-1- SIMILARITY: TO OTHER HERPESVIRUSES GLYCOPROTEIN C.					
CC	CC -1- SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN GENE SUPERFAMILY.				
CC	CC This SWISS-PROT entry is copyrighted. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation of the European Bioinformatics Institute. There are no restrictions on use by non-profit institutions as long as its content is in no modified and this statement is not removed. Usage by and for commercial entities requires a license agreement. (See <a href="http://www.isb-sib.ch/announce">http://www.isb-sib.ch/announce</a> or send an email to license@isb-sib.ch).				
CC	CC DR PIR: A26097; AAA47464.1; -.				
CC	DR PIR: A26097; VGBEPB.				
KW	Glycoprotein; Transmembrane; Signal.				
FT	SIGNAL 1 22				
FT	CHAIN 23 479				
FT	CARBONYD 40 40				
FT	CARBONYD 84 84				
FT	CARBONYD 169 169				
FT	CARBONYD 192 192				
FT	CARBONYD 220 220				
FT	CARBONYD 228 228				
FT	CARBONYD 285 285				
FT	CARBONYD 302 302				
SQ	SEQUENCE 479 AA; 51206 MW; 42E5703 CRC32;				
Query Match	75.8%; Score 47; DB 1; Length 479;				
Best Local Similarity	75.0%; Pred. No. 5.22e+00;				
Matches	6; Conservative 2; Mismatches 0; Indels 0; Gaps 0				
Db	456 AGIGIIAI 463				
Qy	: 3 AGIGIIIV 10				
RESULT	7	METE_ECOLI	STANDARD;	PRT;	752 AA.
ID	P25665;				
AC	01-MAY-1992 (Rel. 22, Created)				
DT	15-JUL-1998 (Rel. 36, Last sequence update)				
DT	15-DEC-1999 (Rel. 39, Last annotation update)				
DE	5-METHYLTHIODYROPIROTRIGONATE--HOMOCYSTEINE METHYLTRANSFERASE				
DE	(EC 2.1.1.14) (METHIONINE SYNTHASE, VITAMIN-B12 INDEPENDENT ISOZYME)				
DE	(COBALAMIN-INDEPENDENT METHIONINE SYNTHASE).				
GN	ME.				
OS	Escherichia coli.				
OC	Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;				
OC	Escherichia.				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RC	STRAIN=K12 / MG1655;				
RX	MEDLINE: 92358234.				
RX	DANIELS D.L., PLUNKETT G. III, BURLAND V.D., BLATTNER F.R.;				
RT	"Analysis of the Escherichia coli genome: DNA sequence of the region from 84.5 to 86.5 minutes."				
RT	Science 257:771-778(1992).				
RT	RL				



CC	-1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
CC	-1- SIMILARITY: BELONGS TO THE UPF0056 (MARC) FAMILY.
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <a href="http://www.isb-sib.ch/announce/">http://www.isb-sib.ch/announce/</a> or send an email to license@isb-sib.ch).
CC	or send an email to license@isb-sib.ch).
CC	DR EMBL; AP000003; BAA20851.1; -.
CC	RW Hypothetical protein; Transmembrane.
AC P36136;	PFT TRANSEM 8 28 POTENTIAL.
DT 01-JUN-1994 (Rel. 29, Created)	PFT TRANSEM 49 69 POTENTIAL.
DT 01-JUN-1994 (Rel. 29, Last sequence update)	PFT TRANSEM 73 93 POTENTIAL.
DT 01-JUN-1994 (Rel. 29, Last annotation update)	PFT TRANSEM 111 131 POTENTIAL.
DE HYPOTHETICAL 31.0 KD PROTEIN IN GAPI-NAP1 INTERGENIC REGION.	PFT TRANSEM 140 160 POTENTIAL.
DF YKR043C.	PFT TRANSEM 181 201 POTENTIAL.
GN OS Saccharomyces cerevisiae (Baker's yeast).	SEQUENCE 201 AA; 21592 MW; 97675186 CRC32;
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccaromycetales; Saccharomycetaceae; Saccharomyces.	Query Match Score 45; DB 1; Length 201; Best Local Similarity 60.0%; Pred. No. 1.46e+01; Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
RN [1]	Db 137 ESPGIVLTI 146
RP SEQUENCE FROM N.A.	Qy 1 EAAGIGILTV 10
RA URBESTARAU L.A.; JAUNITAUX J.-C.;	RESULT 11 Y4TQ_RHISN STANDARD; PRT; 291 AA.
RL Submitted (MAR-1994) to the EMBL/GenBank/DDBJ databases.	ID Q53129; 01-NOV-1997 (Rel. 35, Created)
CC	DT 01-NOV-1997 (Rel. 35, Last sequence update)
CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license@isb-sib.ch/announce/ or send an email to license@isb-sib.ch).	DE PROBABLE PEPTIDE ABC TRANSPORTER PERMEASE PROTEIN Y4TQ.
CC	GN Y4TQ.
DR CAA82119.1; -.	OS Rhizobium sp. (strain NGR334).
DR S38115; S38115.	OG Plasmid sym PNG234a.
DR PFAM; PF00300; PGAM; 1.	CC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group; Rhizobiaceae; Rhizobium.
KW HYPOTHETICAL PROTEIN.	RN [1]
SQ SEQUENCE 271 AA; 31022 MW; F8A036AB CRC32;	RP SEQUENCE FROM N.A.
Query Match Score 46; DB 1; Length 271; Best Local Similarity 55.6%; Pred. No. 8.7e+00; Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;	RA MEDLINE; 97305936.
Db 233 DAGGIGVL 241	RA FREIBERG C.A., FELLAY R., BAIROCH A., BROUGHTON W.J., ROSENTHAL A., PERRET X.;
Qy 1 EAAGIGILT 9	RT "Molecular basis of symbiosis between Rhizobium and legumes." ;
RESULT 10 Y760_PYRHO STANDARD; PRT; 201 AA.	RT Nature 387:394-401(1997).
ID Y760_PYRHO	RN [2]
AC O38499;	RP SEQUENCE FROM N.A.
DT 15-DEC-1998 (Rel. 37, Created)	RX MEDLINE; 96389014.
DT 15-DEC-1998 (Rel. 37, Last sequence update)	RA FREIBERG C., PERRET X., BROUGHTON W.J., ROSENTHAL A.;
DT 15-DEC-1999 (Rel. 39, Last annotation update)	RT "Sequencing the 500-kb GC-rich symbiotic replicon of Rhizobium sp. NGR234 using dye terminators and a thermostable 'sequenase'; a beginning." ;
DE PH0760 OR PHC1026.	RT Genome Res. 6:590-600(1996).
OS Pyrococcus horikoshii.	CC -1- FUNCTION: PROBABLY PART OF A BINDING-PROTEIN-DEPENDENT TRANSPORT SYSTEM Y4TQOPRS FOR A PEPTIDE. PROBABLY RESPONSIBLE FOR THE TRANSLLOCATION OF THE SUBSTRATE ACROSS THE MEMBRANE.
RN [1]	CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. INNER MEMBRANE (POTENTIAL).
RP SEQUENCE FROM N.A.	CC -1- SIMILARITY: WITH INTEGRAL MEMBRANE COMPONENTS OF OTHER BINDING-PROTEIN-DEPENDENT TRANSPORT SYSTEMS. BELONGS TO THE OPPBC SUBFAMILY.
RC SPTRAIN=OT3;	CC
RX MEDLINE; 98344137.	CC
RA KAWABAYASI Y., SAWADA M., HORIKAWA H., HAICAWA Y., HINO Y., NAGAI Y., YAMAMOTO S., SEKINE M., BABU S., KOSUGI H., OGURA K., TAKAMYA M., OHFUKU Y., SAKAI M., OGURA K., OTSUKA R., NAKAZAWA H., TANAKA T., KUDOH Y., YAMAZAKI J., KUSHIDA N., OGUCHI A., FUNAHASHI T., KIKUCHI H., NAKAMURA Y., ROBB T.F., HORIKOSHI K., MASUCHI Y., SHIZUYA H., KIKUCHI H.;	CC
RA "Complete sequence and gene organization of the genome of a hyperthermophilic archaeon, Pyrococcus horikoshii OT3." ;	CC
RL DNA Res. 5:55-76(1998).	CC

CC | :||||| |  
DR EMBL; Z68203; CAA92399.1;  
DR EMBL; AE00098; AAB91870.1;  
DR PROTE; PS00402; BPD\_TRANSP\_INN\_MEMBER; 1.  
DR PFAM; PF00528; BPD\_transp; 1.  
KW Hypothetical protein; Transport; Amino-acid transport; Transmembrane;  
KW • Inner membrane; PlasmaM.  
FT TRANSEM 28 48 POTENTIAL.  
FT TRANSEM 92 112 POTENTIAL.  
FT TRANSEM 137 157 POTENTIAL.  
FT TRANSEM 213 233 POTENTIAL.  
FT TRANSEM 249 269 POTENTIAL.  
SQ SEQUENCE 291 AA; 30910 MW; 3263271E CRC32;

Query Match 72.6%; Score 45; DB 1; Length 291;  
Best Local Similarity 66.7%; Pred. No. 1.46e+01;  
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
RN [1]

Db 147 GPGIGILIV 155  
:||||| |  
QY 2 AAGIGILTV 10

RESULT 12  
ID FTSZ\_AZ0V1 STANDARD PRT; 394 AA.  
AC P77B17;  
DT 01-NOV-1997 (Rel. 35, Created)  
DT 15-DEC-1999 (Rel. 39, Last sequence update)  
DE CELL DIVISION PROTEIN FTSZ.  
RA Azotobacter vinelandii.  
OS Azotobacter vinelandii.  
OC Bacteria; Proteobacteria; gamma subdivision; Azotobacteraceae;  
OC Azotobacter.  
RN [1]  
RP SEQUENCE FROM N.A., AND CHARACTERIZATION.  
RC STRAIN=DJ116;  
RX MEDLINE: 98267010.  
RA LU C., STRICKER J., ERICKSON H.P.;  
RT maritima -- quantitation, GTP hydrolysis, and assembly.;  
RL Cell Motil. Cytoskeleton 40:71-86(1998).  
RA -- FUNCTION: THIS PROTEIN IS ESSENTIAL TO THE CELL-DIVISION PROCESS.  
CC ITS SEEMS TO ASSEMBLE INTO A DYNAMIC RING ON THE INNER SURFACE OF  
THE CITOPLASMIC MEMBRANE AT THE PLACE WHERE DIVISION WILL OCCUR,  
CC BEGIN. BINDS TO AND HYDROLYSES GTP.  
CC -1- SUBUNIT: AGGREGATE TO FORM A RING-LIKE STRUCTURE.  
CC -1- SUBCELLULAR LOCATION: CITOPLASMIC. ASSEMBLE AT THE INNER SURFACE  
OF THE CITOPLASMIC MEMBRANE (BY SIMILARITY).  
CC -1- SIMILARITY: BELONGS TO THE FTSZ FAMILY.

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CC DR AAC24603; U65939; AAC25784; 1;  
DR HSSP; Q57816; 1ESZ.  
DR PROTE; PS01134; FTSZ-1; 1.  
DR PROTE; PS01135; FTSZ-2; 1.  
KW Cell division; Septation; GTP-binding.  
FT NP\_BIND 104 112 GTP (POTENTIAL).  
SQ SEQUENCE 394 AA; 4E887134 CRC32;

Query Match 71.0%; Score 44; DB 1; Length 394;  
Best Local Similarity 77.8%; Pred. No. 2.41e+01;  
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Db 121 AKGLGLTV 129

QY 1 :||||| |  
2 AAGIGILTV 10

RESULT 13  
ID GLMU\_ECOLI STANDARD PRT; 456 AA.  
AC P77114; P76746;  
DT 01-AUG-1990 (Rel. 15, Created)  
DT 01-NOV-1997 (Rel. 35, Last sequence update)  
DT 01-NOV-1997 (Rel. 35, Last annotation update)  
DE UDP-N-ACETYLGLUCOSAMINE PYROPHOSPHORYLASE (EC 2.7.7.23) (N-  
DE ACETYLGLUCOSAMINE-1-PHOSPHATE URIDYLTRANSFERASE).  
GN GLMU.  
OS Escherichia coli.  
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;  
OC Escherichia.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE: 85121806.  
RA WALKER J.E., GAY N.J., SARASSE M., EBERLE A.N.;  
RT "DNA sequence around the Escherichia coli unc operon. Completion of  
the sequence of a 17 kilobase segment containing astA, oriC, unc,  
glnS and phoS.";  
RT Biochem. J. 224:799-815(1984).  
RN [2]  
RP SEQUENCE FROM N.A.  
RC STRAIN=K12 / MG1655;  
RX MEDLINE: 93315143.  
RA BURLAND V.D., PLUNKETT G. III, DANIELS D.L., BLATTNER F.R.;  
RT "DNA sequence and analysis of 136 kilobases of the Escherichia coli  
genome: organizational symmetry around the origin of replication.";  
RL Genomics 16:551-561(1993).  
RN [3]  
RP REVISIONS.  
RC STRAIN=K12 / MG1655;  
RX MEDLINE: 97426617.  
RA BLATTNER F.R., PLUNKETT G. III, BLOCH C.A., PERNA N.T., BURLAND V.,  
RA RILEY M., COLLADO-VIDES J., GLASNER J.D., RODE C.K., MAYHEW G.F.,  
RA GREGOR J., DAVIS N.W., KIRKPATRICK H.A., GOEDEN M.A., ROSE D.J.,  
RA MAU B., SHAO Y.;  
RT "The complete genome sequence of Escherichia coli K-12.";  
RL Science 277:1453-1474(1997).  
RN [4]  
RP IDENTIFICATION.  
RX MEDLINE: 94012475.  
RA MENGIN-LERICREUX D., VAN HELLENOORT J.;  
RT "Identification of the glmU gene encoding N-acetylglucosamine-1-  
phosphate uridylyltransferase in Escherichia coli.";  
RL J. Bacteriol. 175:6150-6157(1993).  
CC FUNCTION: BIFUNCTIONAL ENZYME RESPONSIBLE FOR THE ACETYLATION OF  
GLC-N-1-P TO GIVE GLCNAC-1-P AND THE SYNTHESIS OF UDP-GLCNAc.  
CC -1- CATALYTIC ACTIVITY: UTP + N-ACETYL-ALPHA-D-GLUCOSAMINE  
CC -1- PHOSPHATE = PYROPHOSPHATE + UDP-N-ACETYL-D-GLUCOSAMINE.  
CC -1- PATHWAY: PEPTIDOLYSACCHARIDE BIOSYNTHESIS.  
CC -1- SIMILARITY: BELONGS TO THE CYSE/LACPA/NODL FAMILY OF  
ACETYLTRANSFERASES. COMPOSED OF MULTIPLE REPEATS OF [LIV]-G-X(4).  
CC -1- CAUTION: REF. 2 SEQUENCE DIFFERS FROM THAT SHOWN DUE TO A  
CC FRAMESHIFT THAT CREATES TWO ORFS.  
CC -1-  
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CC DR EMBL; X01631; CAA25784; 1;  
CC DR EMBL; L10328; AAC2082; 1; ALT\_FRAME.  
CC DR EMBL; L10328; AAA2081; 1; ALT\_FRAME.  
CC DR AE000450; AAC76753; 1; -.  
CC DR ECOGENE; EG11198; GLMU  
CC PROSITE; PS00101; HEXADEP\_TRANSFERASE; 1.

DR PFAM; PF00132; hexapep; 3 .  
 DR PFAM; PF00483; NTP transferase; 1 ; Transferase;  
 KW Peptidoglycan synthesis; Cell wall; Transferase;  
 KW Nucleotidyltransferase; Repeat; Multifunctional enzyme.  
 FT CONFLICT 186 187 KL->R (IN REF. 1).  
 SQ SEQUENCE 456 AA; 49190 MW; B9E65439 CRC32;

Query Match 71.0%; Score 44 ; Length 456;  
 Best Local Similarity 75.0%; Pred. No. 2.41e+01;  
 Matches 6; Conservative 2; Mismatches 0; Gaps 0; Indels 0; Gaps 0;

Db 124 GIGILTV 131  
 QY :|||:|||  
 QY 3 AGIGILTV 10

RESULT 14  
 ID FLID\_ECOLI STANDARD; PRT; 467 AA.  
 AC P2416;  
 DT 01-MAR-1992 (Rel. 21, Created)  
 DT 01-AUG-1992 (Rel. 23, Last sequence update)  
 DT 01-NOV-1997 (Rel. 35, Last annotation update)  
 DE FLAGELLAR HOOK-ASSOCIATED PROTEIN 2 (HAP2) (FILAMENT CAP PROTEIN).  
 GN FLID OR FLBC OR FLAV.  
 OS Escherichia coli.  
 OC Bacteriia; Proteobacteria; gamma subdivision; Enterobacteriaceae;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=J41;  
 RX MEDLINE; 92407478.  
 RA KAWAGISHI I., MUELLER V., WILLIAMS A.W., IRIKURA V.M., MACNAB R.M.;  
 RT "Subdivision of flagellar region III of the Escherichia coli and  
 salmonella typhimurium chromosomes and identification of two  
 additional flagellar genes.";  
 RL J. Gen. Microbiol. 138:1065(1992).  
 [2]  
 RN SEQUENCE FROM N.A.  
 RC STRAIN=K12 / MG1652;  
 RX MEDLINE; 97426617.  
 RA BLATTNER F.R., PLUNKETT G. III, BLOCH C.A., PERNA N.T., BURLAND V.,  
 RA RILEY M., COLLADO-VIDES J., GLASNER J.F., RODE C.K., MAYHEW G.F.,  
 RA GREGOR J., DAVIS N.W., KIRKPATRICK H.A., GOEDEN M.A., ROSE D.J.,  
 RA MAU B., SHAO Y.;  
 RT "The complete genome sequence of Escherichia coli K-12.";  
 RL Science 277:1453-1474 (1997).  
 [3]  
 RN SEQUENCE OF 1-152 FROM N.A.  
 RC STRAIN=K12;  
 RX MEDLINE; 89281489.  
 RA HANAFUSA T., SAKAI A., TOMINAGA A., ENOMOTO M.;  
 RT "Isolation and characterization of Escherichia coli hag operator  
 mutants whose hag4 expression has become repressible by a Salmonella  
 H1 repressor.";  
 RL Mol. Gen. Genet. 216:44-50 (1989).  
 [4]  
 RN SEQUENCE OF 1-8 FROM N.A.  
 RX MEDLINE; 83238225.  
 RA SZKELELY E., SIMON M.;  
 RT "DNA sequence adjacent to flagellar genes and evolution of flagellar-  
 phase variation.";  
 RL J. Bacteriol. 155:74-81(1983).  
 CC -I- FUNCTION: CAPPING PROTEIN FOR THE FLAGELLA; FORMS THE DISTAL END  
 CC -I- SIMILARITY: TO OTHER FILAMENT CAP PROTEINS.  
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CC EMBL; M85240; AAC23790.1; -.  
 DR EMBL; AE000385; AAC7491.1; -.  
 DR EMBL; X17440; CAA35487.1; -.  
 DR EMBL; J01607; AAC92490.1; -.  
 DR PIR; PV0005; PV0005.  
 DR ECOGENE; EG10841; FLID.  
 RW Flagella.

FT INIT MET 0 0 BY SIMILARITY.  
 FT CONFFLICT 113 113 T->R (IN REF. 3).  
 SQ SEQUENCE 467 AA; 48270 MW; 14800AE CRC32;

Query Match 71.0%; Score 44 ; DB 1; Length 467;  
 Best Local Similarity 50.0%; Pred. No. 2.41e+01;  
 Matches 5; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Db 368 DASGVGALIV 377  
 ID XYND\_PAEPO STANDARD; PRT; 635 AA.  
 AC P45766;  
 DT 01-NOV-1995 (Rel. 32, Created)  
 DT 01-NOV-1995 (Rel. 32, Last sequence update)  
 DT 15-JUL-1999 (Rel. 38, Last annotation update)  
 DE ENDO-1,4-BETA-XYLANASE D PRECURSOR (EC 3.2.1.8) (XYLANASE D).  
 GN XND.  
 OS Paenibacillus polymyxa (Bacillus polymyxa).  
 OC Bacteria; Firmicutes; Bacillales/Clostridium group;  
 OC Beccilli/Stephyllococcus group; Paenibacillus.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=ATCC 842;  
 RX MEDLINE; 92041687.  
 RA GOSALBES M.J., PIREZ-GONZALEZ J.A., GONZALEZ R., NAVARRO A.;  
 RT "Two beta-glycanase genes are clustered in *Bacillus polymyxa*:  
 RT molecular cloning, expression, and sequence analysis of genes  
 RT encoding a xylanase and an endo-beta-(1,3)-(1,4)-glucanase.";  
 RL J. Bacteriol. 173:705-7710(1991).  
 CC -I- FUNCTION: SHOWS XYLANASE ACTIVITY AS WELL AS ALPHA-L-  
 CC ARABINOFRURANOSIDASE ACTIVITY.  
 CC -I- CATALYTIC ACTIVITY: ENDOHYDROLYSIS OF 1,4-BETA-D-XYLOSIDIC  
 CC LINKAGES IN XYLANS.  
 CC -I- PATHWAY: XYLAN DEGRADATION.  
 CC -I- SIMILARITY: BELONGS TO FAMILY 43 OF GLYCOSYL HYDROLASES.

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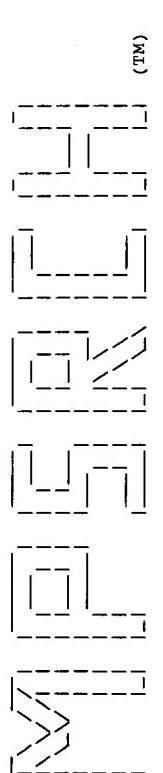
CC DR EMBL; X57034; CAA40378.1; -.  
 CC KW Xylan degradation; Hydrolase; Glycosidase; Signal.  
 CC FT SIGNAL 1 26 POTENTIAL.  
 CC FT CHAIN 27 635 ENDO-1,4-BETA-XYLANASE D.  
 CC SQ SEQUENCE 635 AA; 67914 MW; 078AA82 CRC32;

Query Match 71.0%; Score 44 ; DB 1; Length 635;  
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 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Db 149 GAGIGVLT 156  
 QY :||||:|||  
 QY 2 AAGIGILTV 9

Search completed: Fri May 5 22:09:57 2000

Job time : 39 secs.

\*\*\*\*\*  
  
\*\*\*\*\*

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MPSrch\_PP protein - protein database search, using Smith-Waterman algorithm

Run on: Fri May 5 22:08:01 2000; MasPar time 4.93 Seconds

95.764 Million cell updates/sec  
Tabular output not generated.

Title: >US-09-267-439-17  
Description: (1-10) from US09267439.pep

Perfect Score: 62  
Sequence: EAAGIGILTV 10

Scoring table: PAM 150  
Gap 15

Searched: 142080 seqs, 47172406 residues

Post-processing: Minimum Match 0%  
Listing first 45 summaries

Database: pir62  
1:pir1 2:pir2 3:pir3 4:pir4

Statistics: Mean 22.809; Variance 26.769; scale 0.852

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No. Score Query Match Length DB ID Description Pred. No.

Result No.	Score	Query	Match	Length	DB	ID	Description	Pred. No.
1	62	100.0	118	2	A55253		melanoma antigen MART	6.18e-03
2	48	77.4	333	1	DEALXE		acetoin[2,6-dichlorop	9.75e+00
3	47	75.8	101	2	A33351		H+-transporting ATP S	1.57e+01
4	47	75.8	110	1	VHBPD1		major capsid protein	1.57e+01
5	47	75.8	165	2	C70959		hypothetical protein	1.57e+01
6	47	75.8	250	2	A69843		hypothetical protein	1.57e+01
7	47	75.8	479	1	YGBBPB		glycoprotein GII pre	1.57e+01
8	47	75.8	753	2	A42863		5-methyltetrahydropo	1.57e+01
9	47	75.8	773	2	T00502		protein kinase homolo	1.57e+01
10	46	74.2	231	1	S48276		YSA1 protein - Yeast	2.52e+01
11	46	74.2	271	2	S38115		hypothetical protein	2.52e+01
12	45	72.6	201	2	A71124		hypothetical protein	4.01e+01
13	45	72.6	339	2	S62369		methylcobalamin-coen	4.01e+01
14	45	72.6	420	2	S29131		Kan-1 protein - rat	4.01e+01
15	45	72.6	744	2	A43353		ascites stiaglycopro	4.01e+01
16	44	71.0	98	2	D72106		hypothetical protein	6.33e+01
17	44	71.0	456	2	C65176		glmu protein - Escher	6.33e+01
18	44	71.0	468	2	A54956		flagellar hook associ	6.33e+01
19	44	71.0	620	2	H69382		ABC transporter, ATP-	6.33e+01
20	44	71.0	635	2	S19011		endo-1,4-beta-xylanas	6.33e+01
21	43	69.4	123	2	A71312		probable anti-sigma F	9.91e+01
22	43	69.4	132	2	S01903		H+-transporting ATP S	9.91e+01
23	43	69.4	207	2	H69694		ribosomal protein L4	9.91e+01

24 43 69.4 231 2 G75193 9.91e+01  
25 43 69.4 345 1 K270602 9.91e+01  
26 43 69.4 402 2 D70602 9.91e+01  
27 43 69.4 461 2 D70073 9.91e+01  
28 43 69.4 461 2 T03561 9.91e+01  
29 43 69.4 493 1 ACMSE 9.91e+01  
30 43 69.4 509 2 H70597 9.91e+01  
31 43 69.4 580 2 F70868 9.91e+01  
32 43 69.4 590 2 A45772 9.91e+01  
33 43 69.4 593 1 A69655 9.91e+01  
34 43 69.4 611 2 JT0592 9.91e+01  
35 43 69.4 667 2 F70682 9.91e+01  
36 43 69.4 675 2 S53832 9.91e+01  
37 43 69.4 729 2 T0617 9.91e+01  
38 43 69.4 746 2 T06017 9.91e+01  
39 43 69.4 1217 1 EGMSMG 9.91e+01  
40 43 69.4 1325 2 A64905 9.91e+01  
41 43 69.4 1436 2 S67655 9.91e+01  
42 43 69.4 1616 2 G70668 9.91e+01  
43 42 67.7 218 1 B41316 1.54e+02  
44 42 67.7 251 2 S56416 1.54e+02  
45 42 67.7 417 2 D70321 1.54e+02

#### ALIGNMENTS

RESULT 1 #type complete  
ENTRY A55253 #type antigen MART-1 - human  
TITLE melanoma antigen MART-1  
ALTERNATE\_NAMES melan A protein  
ORGANISM Homo sapiens #common\_name man  
DATE 06-Feb-1995 #sequence\_revision 06-Feb-1995 #text\_change  
10-Sep-1997  
ACCESSIONS A55253; I38506  
REFERENCE A55253  
#authors Kawakami, Y.; Eliyahu, S.; Delgado, C.H.; Robbins, P.F.; Rivoltini, L.; Topalian, S.L.; Maki, T.; Rosenberg, S.A.  
#journal Proc. Natl. Acad. Sci. U.S.A. (1994) 91:1515-1519  
#title Cloning of the gene coding for a shared human melanoma antigen recognized by autologous cytolytic T cells infiltrating into tumor.  
#cross-references MUID:94224770  
#accesion A55253  
#status preliminary  
##molecule-type mRNA  
##residues 1-118 ##label RAW  
REFERENCE I38506  
#authors Couille, P.G.; Brichard, V.; Van Pel, A.; Wolfel, T.; Schneider, J.; Traversari, C.; Mattei, S.; De Paepe, E.; Lurquin, C.; Szilvai, J.P.; Renaud, J.; Boon, T.  
#journal J. Exp. Med. (1994) 180:35-42  
#title A new gene coding for a differentiation antigen recognized by comments:  
#cross-references MUID:94275389  
#accesion I38506  
#status preliminary; translated from GB/EMBL/DDJB  
##molecule-type mRNA  
##residues 1-118 ##label RES  
SUMMARY #length 118  
GENETICS #cross-references EMBL:U06654; NID:9517022; PID:9517023  
#gene GDB:MIANA  
##cross-references GDB:358979  
#map\_position 17q21-17q24  
##molecule-type mRNA  
##residues 1-118 #label RES  
#length 118  
Query Match 100.0%; Score 62; DB 2; Length 118;  
Best Local Similarity 100.0%; Pred. No. 6.18e-03;  
Matches 0; Mismatches 0; Indels 0; Gaps 0;

Db 26 EAAGIGILTV 35  
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Qy 1 EAAGIGILTV 10
Qy 1 EAAGIGILTV 9
Qy |||||:||| 10
Qy 2 AAAGIGILTV 9
Qy |||||:||| 9

RESULT 4
ENTRY VHBPDLL #type complete
TITLE major capsid protein D - phage lambda
ORGANISM #head protein D
DATE #formal_name phage lambda
        13-Jun-1983 #sequence_revision 13-Jun-1983 #text_change
        23-Jul-1999
ACCESSIONS G04333; C43013; A04334; A23206
REFERENCE A94614
#authors Daniels, D.
#title submitted to the Nucleic Acid Sequence Database, September
#cross-references MUID:91286190
#accession B42462
#molecule_type DNA
#residues 1-33 #label PRI
#cross-references GB:M66060; PID:9141892; PIDN:AAA21948_1; PID:9141894
COMMENT This is a component of the enzyme complex that catalyzes 2', 6-dichlorophenolindophenol-dependent cleavage of acetoin into acetate and acetaldehyde. The functional enzyme is a tetramer of two alpha and two beta chains.
CLASSIFICATION #domain thiamine pyrophosphate-binding domain homology
KEYWORDS thiamine pyrophosphate-binding domain homology
FEATURE 145-194
#domain thiamine pyrophosphate-binding domain homology
#superfamily pyruvate dehydrogenase (lipoamide) alpha chain;
#heterotetramer; oxidoreductase
SUMMARY #length 333 #molecular_weight 35375 #checksum 2647
Query Match Score 48; DB 1; Length 333;
Best Local Similarity 87.3%; Pred. No. 9.75e+00;
Matches 7; Conservative 0; Indels 0; Gaps 0;
Db 56 EAAGVGIL 63
||||:||| 1 EAAGIGIL 8
Qy 1 EAAGIGILTV 8

RESULT 3
ENTRY A33351 #type complete
TITLE H+-transporting ATP synthase (EC 3.6.1.34) proteolipid chain
ORGANISM - Sulfolobus acidocaldarius
DATE #formal_name Sulfolobus acidocaldarius
        20-Dec-1989 #sequence_revision 20-Dec-1989 #text_change
        22-Jun-1999
ACCESSIONS A33351
#cross-references MUID:89214142
#status preliminary
#molecule_type DNA
#residues 1-101 ##label DEN
#cross-references GB:J04740; PIDN:AAA72703_1; PID:9152925
CLASSIFICATION #superfamily H+-transporting ATP synthase lipid-binding
KEYWORD hydrolase
SUMMARY #length 101 #molecular_weight 10362 #checksum 4300
Query Match Score 47; DB 2; Length 101;
Best Local Similarity 87.5%; Pred. No. 1.57e+01;
Matches 7; Conservative 0; Indels 0; Gaps 0;
Db 59 AAAGVGLAV 64
||||:||| 1 EAAGIGILTV 10
Qy 1 EAAGIGILTV 10

RESULT 5
ENTRY C70959 #type complete
TITLE hypothetical protein Rv1382 - Mycobacterium tuberculosis
        (strain H37RV)
ORGANISM #formal_name Mycobacterium tuberculosis
DATE 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change
        26-Aug-1999
ACCESSIONS C70959
REFERENCE A70500
#authors Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher,
        C.; Harris, D.; Gordon, S.V.; Eiglemer, K.; Gas, S.; Barry,
        C.E.; Tekwani, F.; Badcock, K.; Basham, D.; Brown, D.;
        Chillingworth, T.; Connor, R.; Davies, R.; Devlin, K. ;
        Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S. ;
        Hornsby, T.; Jagels, K.; Krugh, A.; McLean, J.; Moule, S. ;
        Murphy, L.; Oliver, S.; Osborne, J.; Quail, M.A. ;
        Rajandream, M.A.; Rogers, J.; Rutte, S.; Seeger, K. ;
        Skelton, S.; Squares, S.; Soares, R.; Sulston, J.E. ;
        Taylor, K.; Whitehead, S.; Barrell, B.G.

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#journal Nature (1998) 393:537-544  
#title Deciphering the biology of Mycobacterium tuberculosis from  
#cross-references MUID:98044033  
#accession A09843  
#status Preliminary; nucleic acid sequence not shown;  
##molecule\_type DNA  
##residues 1-250 ##label KUN  
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#cross-references GB:Z81011; GB:AL123456; NID:q3242274; PID:e275153;  
#cross-references GB:9621264  
#experimental\_source strain H37Rv

GENETICS  
#gene Rv1382  
#classification #superfamily Mycobacterium tuberculosis hypothetical protein  
#cross-references Rv1382  
#length 165 #molecular-weight 18189 #checksum 5780

SUMMARY  
Query Match Score 47; DB 2; Length 250;  
Best Local Similarity 75.8%; Pred. No. 1 57e-01;  
Matches 1; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Db 128 AGIGLAI 135  
| |||||:  
Qy 3 AGIGILTV 10

RESULT 6  
ENTRY A69843  
TITLE #type complete  
#hypothetical protein Yba - Bacillus subtilis  
#formal\_name pacillus subtilis  
ORGANISM 05-Dec-1997 #sequence\_revision 05-Dec-1997 #text\_change  
DATE 24-Sep-1998

ACCESSIONS  
A69843  
REFERENCE Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Bertero, M.G.; Bessieres, P.; Bolotin, A.; Borchart, S.; Boris, R.; Boursier, L.; Brans, A.; Braun, M.; Brignell, S.C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Choi, S.K.; Codani, J.J.; Connerton, I.F.; Cummings, N.J.; Daniel, R.A.; Deniot, F.; Devine, K.M.; Duisterhoeft, A.; Ehrlich, S.D.; Emmerson, P.T.; Entian, K.D.; Errington, J.; Fabret, C.; Ferrari, E.; Foulger, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Galizzi, A.; Galleron, N.; Ghim, S.Y.; Glaser, P.; Goffeau, A.; Golightly, E.J.; Grandi, G.; Giuseppi, G.; Guy, B.J.; Haga, K.; Haiech, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Hollsappel, S.; Hosono, S.; Hullio, M.P.; Itaya, M.; Jones, D.; Joris, B.; Karamoto, D.; Kasahara, Y.; Klaerr-Blanchard, M.; Klein, C.; Kobayashi, Y.; Koetter, P.; Konigstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois, S.; Lauber, J.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Maeluer, C.; Medigue, C.; Medina, N.; Mellado, R.P.; Mizuno, M.; Moestl, D.; Nakai, S.; Noback, M.; Noone, D.; O'Reilly, M.; Ogawa, K.; Ogiwara, A.; Oudega, B.; Park, S.H.; Parro, V.; Pohl, T.M.; Portelle, D.; Porwolik, S.; Prescott, G.; Rey, M.; Reynolds, S.; Rose, M.; Sadale, Y.; Sato, T.; Scanlon, E.; Roche, B.; Schleicher, S.; Schroeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Seror, S.J.; Serrac P.; Shin, B.S.; Soldo, B.; Sorokin, A.; Taccioni, E.; Takeuchi, T.; Takashita, H.; Takemaru, K.; Takeuchi, M.; Tamakoshi, A.; Tanaka, T.; Terpsira, P.; Tognoni, A.; Tosato, V.; Uchiyama, S.; Vandembol, M.; Vannier, F.; Vassarotti, A.; Viari, A.; Wambutt, R.; Wedler, E.; Wedler, H.; Weitzneger, T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yata, K.; Yoshida, K.; Yoshihikawa, H.F.; Zumstein, E.; Yoshihikawa, H.; Danchin, A.

#journal Nature (1997) 390:249-256  
#title The complete genome sequence of the Gram-positive bacterium Bacillus subtilis.

#cross-references MUID:98044033  
#accession A09843  
#status Preliminary; nucleic acid sequence not shown;  
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##cross-references GB:Z99110; GB:AL009126; NID:92633472; PID:e1183161;  
#cross-references GB:Z81011; GB:AL123456; NID:q3242274; PID:e275153;  
#experimental\_source strain H37Rv

GENETICS  
#gene Yjba  
#length 250 #molecular-weight 30119 #checksum 5271

SUMMARY  
Query Match Score 47; DB 2; Length 250;  
Best Local Similarity 75.8%; Pred. No. 1 57e-01;  
Matches 1; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Db 89 TDGIGILAV 97  
Qy :|||||:  
2 AAGIGILTV 10

RESULT 7  
ENTRY VGBEPB #type complete  
#glycoprotein gIII precursor - suid herpesvirus 1  
#formal\_name suid herpesvirus 1  
TITLE #sequence\_revision 30-Sep-1987 #text\_change  
ORGANISM 16-Jul-1999  
DATE 30-Sep-1987  
ACCESSIONS  
A26097  
REFERENCE A26097  
#authors Robbins, A.K.; Watson, R.J.; Whealy, M.E.; Hays, W.W.; Enquist, L.W.  
#journal J. Virol. (1986) 58:339-347  
#title Characterization of a Pseudorabies virus glycoprotein gene with homology to herpes simplex virus type 1 and type 2 glycoprotein C.  
#cross-references MUID:86200375  
#accession A26097  
##molecule\_type DNA  
##residues 1-479 ##label ROB  
##cross-references GB:M12778; NID:g334049; PIDN:AAA47464.1; PID:g334050  
#cross-references GB:M12778; NID:g334049; PIDN:AAA47464.1; PID:g334050  
#experimental\_source strain Becker  
CLASSIFICATION #superfamily herpesvirus glycoprotein F  
KEYWORDS glycoprotein F  
FEATURE #domain signal sequence #status predicted #label SIG  
1-22 #product glycoprotein III #status predicted #label GPG  
23-479 #binding\_site carbohydrate (Asn) (covalent)  
#predicted  
#status Predicted  
#predicted  
#length 479 #molecular-weight 51206 #checksum 1630

SUMMARY  
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Best Local Similarity 75.8%; Pred. No. 1.57e-01;  
Matches 2; Conservative 6; Mismatches 0; Indels 0; Gaps 0;

Db 456 AGIGILAI 463  
Qy :|||||:  
3 AGIGILTV 10

RESULT 8  
ENTRY A42863 #type complete  
#5-methyltetrahydropteroyltriglutamate--homocysteine  
#S-methyltransferase (EC 2.1.1.14) - Escherichia coli  
#strain K-12  
TITLE  
#organism Escherichia coli  
#flogenome Escherichia coli  
#date 17-Feb-1994 #sequence\_revision 10-Oct-1997 #text\_change  
#accessions F65187; A42863; S30719; 179560  
#reference A64720



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#cross-references MUID:92327848
#accession S23364
#molecule_type DNA
##residues 1-47 ##label MAW
##cross-references EMBL:X66247; NID:93548; PID:g3549
GENETICS
  #gene SGD:YSA1
    ##cross-references SGD:S0000315; MIPS:YBR111c
      #map_position 2R
        CLASSIFICATION #superfamily YffH protein: mutt domain homology
        FEATURE #domain mutt domain homology #label MUTT
        SUMMARY #length 231 #molecular_weight 26687 #checksum 4809
          Query Match Score 46; DB 1; Length 231;
          Best Local Similarity 74.2%; Pred. No. 2.52e+01;
          Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
        Db 79 GIGILTI 85
        Qy 4 GIGILTY 10

        RESULT 11
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        TITLE hypothetical protein YKR043C - yeast (Saccharomyces cerevisiae)
        ORGANISM #formal_name Saccharomyces cerevisiae
        DATE 03-May-1994 #sequence_revision 03-May-1994 #text_change
        14-Nov-1997
        ACCESSIONS S38115
        REFERENCE S38097
        #authors Urrestarazu, L.A.; Jauniaux, J.C.
        #submission submitted to the Protein Sequence Database, March 1994
        #accession S38115
        #molecule_type DNA
        ##residues 1-271 ##label URR
        ##cross-references EMBL:Z28266; NID:9486490; PID:9486491; MIPS:YKR043C
        ##experimental_source strain S288C
        GENETICS
        #map_position 11R
        SUMMARY #length 271 #molecular_weight 31022 #checksum 8533
          Query Match Score 46; DB 2; Length 271;
          Best Local Similarity 74.2%; Pred. No. 2.52e+01;
          Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
        Db 233 DAGGIGVLS 241
        Qy 1 EAAGIGILT 9

        RESULT 12
        ENTRY A71124 #type complete
        TITLE hypothetical protein PH0760 - Pyrococcus horikoshii
        ORGANISM #formal_name Pyrococcus horikoshii
        DATE 14-Aug-1998 #sequence_revision 14-Aug-1998 #text_change
        ACCESSIONS A71124
        #authors
        Kawarabayasi, Y.; Sawada, M.; Horikawa, H.; Haikawa, Y.; Hino, Y.; Yamamoto, S.; Seine, M.; Baba, S.; Kosugi, H.; Hosoyama, A.; Nagai, Y.; Sakai, M.; Ogura, K.; Otsuka, R.; Nakazawa, H.; Takamaya, M.; Ohfuki, Y.; Funahashi, T.; Tanaka, T.; Kudo, Y.; Yamazaki, J.; Kushida, N.; Ouchi, A.; Aoki, K.; Yoshizawa, T.; Nakamura, Y.; Robb, F.T.; Horikoshi, K.; Masuchi, Y.; Shizuya, H.; Kikuchi, H.
        #journal DNA Res. (1998) 5:55-76
        #title Complete sequence and gene organization of the genome of a hyper-thermophilic archaeabacterium, Pyrococcus horikoshii
        #cross-references MUID:98344137
        #accession A71124

#status preliminary; nucleic acid sequence not shown;
#molecule_type DNA
##residues 1-201 ##label KAW
##cross-references GB:AP000003; NID:93236130; PID:d1030794; PID:g3257168
#experimental_source strain OR3
##note this accession replaces an interim accession for a sequence replaced by GenBank
GENETICS
  #gene PH0760
    CLASSIFICATION #superfamily conserved hypothetical protein MJ1677
    SUMMARY #length 201 #molecular_weight 21592 #checksum 3142
      Query Match Score 45; DB 2; Length 201;
      Best Local Similarity 60.0%; Pred. No. 4.0le+01;
      Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
      Db 137 ESPGIVLT 146
      Qy 1 EAAGIGILTV 10

      RESULT 13
      ENTRY S62369 #type complete
      TITLE methylcobalamin-coenzyme M methyltransferase II -
      Methanosaarcina Barkeri
      #formal_name Methanosaarcina Barkeri
      DATE 19-Mar-1997 #sequence_revision 25-Apr-1997 #text_change
      17-Mar-1999
      ACCESSIONS S62369
      REFERENCE S62368
      #authors Harms, U.; Thauer, R.K.
      #journal Eur. J. Biochem. (1996) 235:653-659
      #title Methylcobalamin-coenzyme M methyltransferase isoenzymes MtAA and MtBa from Methanosaarcina Barkeri: cloning, sequencing and differential transcription of the encoding genes, and functional overexpression of the mtAA gene in Escherichia coli.
      #cross-references MUID:96184544
      #accession S62369
      #status preliminary; nucleic acid sequence not shown
      #molecule_type DNA
      ##residues 1-339 ##label HAR
      #cross-references EMBL:X91894; NID:91107727; PID:91107728
      SUMMARY #length 339 #molecular_weight 36761 #checksum 6431
        Query Match Score 45; DB 2; Length 339;
        Best Local Similarity 75.0%; Pred. No. 4.0le+01;
        Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
        Db 307 AGVGLLT 314
        Qy 3 AGIGILTV 10

      RESULT 14
      ENTRY S59131 #type complete
      TITLE Kan-1 protein - rat
      ORGANISM #formal_name Rattus norvegicus #common_name Norway rat
      DATE 15-Feb-1996 #sequence_revision 01-Mar-1996 #text_change
      07-May-1999
      ACCESSIONS S59131
      REFERENCE
      #authors Furutani, M.; Arii, S.; Higashitsuji, H.; Mise, M.; Fukumoto, M.; Takano, S.; Nakayama, H.; Imamura, M.; Fujita, J.
      #journal Biochem. J. (1995) 311:203-208
      #title Reduced expression of kan-1 (encoding putative bile acid-CoA amino acid N-acyltransferase) mRNA in livers of rats after partial hepatectomy and during sepsis.
      #cross-references MUID:96003917
      #accession S59131
      #molecule_type mRNA

```

```

##residues 1-420 ##label FUR
##cross-references EMBL:NID:g604901; PID:q1008487; PID:g604902
SUMMARY #length 420 #molecular_weight 46196 #checksum 4868
* Best Local Similarity 72.6%; Score 45; DB 2; Length 420;
  Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
Db 226 GPGVGILSV 234
Qy 2 AAGIGILTV 10
          ::|||:|||:|
          Qy 2 AAGIGILTV 10

RESULT 15
ENTRY A43353 #type fragment
TITLE ascites sialoglycoprotein-2 - rat (fragment)
ORGANISM #formal_name Rattus norvegicus #common_name Norway rat
DATE 10-Jun-1993 #sequence_revision 18-Nov-1994 #text_change
16-Jul-1999
ACCESSIONS A43353
REFERENCE Sheng, Z.; Wu, K.; Carraway, K.L.; Fregien, N.
#authors J. Biol. Chem. (1992) 267:16341-16346
#title Molecular cloning of the transmembrane component of the 13762
#note mammary adenocarcinoma sialomucin complex. A new member of
the epidermal growth factor superfamily.
#cross-references MUID:92355597
#accession A43353
##status preliminary
##molecule_type mRNA; protein
##residues 1-744 ##label SHE
##experimental_source mammary adenocarcinoma
##note sequence extracted from NCBI backbone (NCBIN:110590,
NCBIP:110691)
CLASSIFICATION #superfamily EGF homology
KEYWORDS glycoprotein
FEATURE 655-694 #domain EGF homology #label EGF
SUMMARY #length 744 #checksum 2462

Query Match 72.6%; Score 45; DB 2; Length 744;
Best Local Similarity 60.0%; Pred. No. 4.0e+01;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
Db 201 ETNGIGLLGV 210
          ::|||:|||:|
          Qy 1 EAAGIGILTV 10

Search completed: Fri May 5 22:09:02 2000
Job time : 61 secs.

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Qy 2 AGIGILTVI 10

RESULT 12 PRELIMINARY; PRT; 677 AA.

ID Q89853; AC Q89853;

DT 01-NOV-1996 (TREMBLrel. 01, Created)

DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)

DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)

DE VIRION SPIKE GLYCOPROTEIN PRECURSOR.

OS Ebola virus (Ebo).

OC Viruses; ssRNA negative-strand viruses; Mononegavirales; Filoviridae;

OC Filovirus.

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=RESTON SUBTYPE, SIENA STRAIN (1992);

RX MEDLINE; 96195018.

RA SANCHEZ A., TRAPPIER S.G., MAHY B.W.J., PETERS C.J., NICHOL S.T.;

RT "The virion glycoproteins of Ebola viruses are encoded in two reading frames and are expressed through transcriptional editing.";

RL Proc. Natl. Acad. Sci. U.S.A. 93:3602-3607(1996).

DR EMBL; U23417; AAC54891.1; -.

DR EMBL; U23416; AAC54899.1; -.

DR PFAM; PF01611; Filo\_glycop; 1.

KW Signal.

FT SIGNAL 1 33 POTENTIAL.

SQ SEQUENCE 677 AA; 74523 MW; C56FB3E0 CRC32;

Query Match 76.2%; Score 48; DB 14; Length 677;

Best Local Similarity 70.0%; Pred. No. 2.20e+01;

Matches 6; Conservative 0; Mismatches 2; Indels 1; Gaps 0;

Db 654 PAGIGILTVI 663

Qy 1 AGIGILTVI 10

RESULT 13 PRELIMINARY; PRT; 677 AA.

ID Q66799; AC Q66799;

DT 01-NOV-1996 (TREMBLrel. 01, Created)

DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)

DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)

DE VIRION SPIKE GLYCOPROTEIN PRECURSOR.

GN GP.

OS Ebola virus (Ebo), and Ebola virus Reston.

OC Viruses; ssRNA negative-strand viruses; Mononegavirales; Filoviridae;

OC Filovirus.

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=RESTON SUBTYPE, RESTON STRAIN;

RX MEDLINE; 96195019.

RA SANCHEZ A., TRAPPIER S.G., MAHY B.W.J., PETERS C.J., NICHOL S.T.;

RT "The virion glycoproteins of Ebola viruses are encoded in two reading frames and are expressed through transcriptional editing.";

RL Proc. Natl. Acad. Sci. U.S.A. 93:3602-3607(1996).

RN [2]

RP SEQUENCE FROM N.A.

RC STRAIN=PENNSYLVANIA;

RX MEDLINE; 98245155.

RA VOLCHKOV V.E., FELDMANN H., VOLCHKOVA V.A., KLENK H.D.;

RT "Processing of the Ebola virus glycoprotein by the protease convertase furin.";

RL Proc. Natl. Acad. Sci. U.S.A. 95:5762-5767(1998).

DR EMBL; U23154; AAC54881.1; -.

DR EMBL; AF034645; AAC24346.1; -.

DR PFAM; PF01611; Filo\_glycop; 1.

KW Signal.

FT SIGNAL 1 33 POTENTIAL.

SQ SEQUENCE 677 AA; 74432 MW; 9EA5B80C CRC32;

Query Match 76.2%; Score 48; DB 14; Length 677;

Best Local Similarity 70.0%; Pred. No. 2.20e+01;

Db 379 SGVGLVYY 387

:1:1:1 :1:

Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0; Search completed: Fri May 5 22:20:06 2000  
 Job time : 89 secs.

---

Db 654 PAGIGLIGVI 663  
 Qy 1 AAGIGILTVI 10

RESULT 14  
 ID Q9ZT37 PRELIMINARY; PRT; 808 AA.  
 AC Q9ZT37;  
 DT 01-MAY-1999 (TRIMBLrel. 10, Created)  
 DT 01-MAY-1999 (TRIMBLrel. 10, Last sequence update)  
 DT 01-MAY-1999 (TRIMBLrel. 10, Last annotation update)  
 DE PUTATIVE GLUTAMATE RECEPTOR.  
 GN GLR1  
 OS Arabidopsis thaliana (Mouse-ear cress).  
 OC Eukarya; Viricoplantae; Streptophyta; Embryophyta; Tracheophyta;  
 CC euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;  
 CC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;  
 CC Arabidopsis.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE: 99039437.  
 RA LAM H.M., CHIU J., HSIEH M.H., MEISEL L., OLIVEIRA I.C., SHIN M.,  
 RA CORUZZI G.;  
 RT "Glutamate receptor genes in plants.";  
 RL Nature 396:125-126 (1998).  
 DR EMBL; AF079998; AAD09173..1.; -.  
 RW Receptor.  
 SQ SEQUENCE 808 AA; 90518 MW; C3554B89 CRC32;

Query Match 76.2%; Score 48; DB 10; Length 808;  
 Best Local Similarity 100.0%; Pred. No. 2.20e+01;  
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 519 GIGILTV 525  
 Qy 3 GIGILTV 9

RESULT 15  
 ID Q49684 PRELIMINARY; PRT; 91 AA.  
 AC Q49684;  
 DT 01-NOV-1996 (TRIMBLrel. 01, Created)  
 DT 01-NOV-1996 (TRIMBLrel. 01, Last sequence update)  
 DT 01-NOV-1996 (TRIMBLrel. 01, Last annotation update)  
 DE B1496\_C2\_163.  
 OS Mycobacterium leprae.  
 CC Bacteria; Firmicutes; Actinobacteria; Actinomycetinae; Mycobacterium.  
 CC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RA ROBISON K.;  
 RL Submitted (NOV-1993) to the EMBL/GenBank/DDBJ databases.  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RA ROBISON K.;  
 RL Submitted (JAN-1994) to the EMBL/GenBank/DDBJ databases.  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RA ROBISON K.;  
 RL Submitted (MAR-1994) to the EMBL/GenBank/DDBJ databases.  
 DR EMBL; U00013; AAA17122.1;  
 SQ SEQUENCE 91 AA; 9561 MW; A1C4ED5D CRC32;

Query Match 74.6%; Score 47; DB 2; Length 91;  
 Best Local Similarity 66.7%; Pred. No. 3.48e+01;  
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Db 56 AGIGVLSAI 64  
 Qy 1 |||||: | 2 AGIGILTVI 10

Release 3.1A John F. Collins, Biocomputing Research Unit.  
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protein - protein database search, using Smith-Waterman algorithm  
run on: Fri May 5 22:17:41 2000; Maspair time 3.08 Seconds  
96,985 Million cell updates/sec  
molecular output not generated.

Prescription:  
perfect Score:  
63  
(1-10) from US09267439 .pep

Sequence: 1 AAGIGIILTVI 10

scoring table: PAM 150  
Gap 15

searched: 82229 seqs, 29864866 residues

post-processing: Minimum Match 0% Listing first 45 summaries

database: swiss-prot38

```

statistics: Mean 23.818; Variance 26.551; scale 0.897
1:swisspiul

```

Pred. No. is the number of results predicted by chance to have a

and is derived by analysis of the total score distribution.

DRAFT - 2000-02-08  
Query result

No.	Score	Match Length	DB ID	Description	Pred.	No.
1	63	100	0	MAB1 UTINAN	METANOMA	3 280-03

2	52	82.5	479	1	VGLC_PRVTF	GLYCOPROTEIN GLII PREC	2.10e-03
3	50	79.4	461	1	YXCC_BACSU	HYPOTHETICAL METABOLIT	2.87e+00
2	52	100.0	479	1	YXCC_BACSU	HYPOTHETICAL METABOLIT	2.87e+00

4	49	77.8	345	1 HRCA_STRMU
5	49	77.8	883	1 YDGH_BACSU
6	48	76.2	231	1 VYRASU
				PUTATIVE MEMBRANE PROT
				VSA1 PROTEIN
				4.74e+00
				4.74e+00
				7.76e+00

9	48	76.2	633	1	Y561-HAETN	HYPOTHETICAL PROTEIN H	7.76e+00
10	48	76.2	1331	1	CYB1-SU1AC	RECEPTOR-TYPE ADENYLAT	7.76e+00
11	47	74.6	101	1	ATPL_SUITAC	MEMBRANE-ASSOCIATED AT	1.26e+01

		SECD_HELPP	HYPOTHETICAL PROTEIN P
14	47	526	1
15	46	73.0	201
16	46	73.0	337
			1 OPSX_HUMAN

17	46	73.0	440	1	UGTC-CAEFL PUTATIVE UDP-GLUCURONO 2.03e+01
18	46	73.0	977	1	YGSB-CHPO HYPOTHETICAL 2.03e+01
19	46	73.0	1100	1	CVCD CANED DOMINANT CHAVIY 2.03e+01

20	45	71.4	1.235	1 YATQ_ECOLI	HYPOTHETICAL
20	45	71.4	2.935	1 YATQ_RHISN	24.6 KDP
21	45	71.4	2.91	1 PROBABLE PEPTIDE ABC	T 3.24e+01

24	44	69..8	27..8	1	YOHABACSU	HYPOTHETICAL	31..8	KD P	5..13e+01
25	44	69..8	28..9	1	HTBLCLOTS	3-HYDROXYBUTYRYL-COA D	5..13e+01		
25	44	69..8	39..4	1	FTISZAZOVI	CELL DIVISION PROTEIN	5..13e+01		
27	44	69..8	45..6	1	GILMUECOLI	UDP-N-ACETYLGALOSAMIN	5..13e+01		
28	44	69..8	61..0	1	FIMBLIDICDI	FIMBRIN	5..13e+01		
29	44	69..8	61..1	1	YD3MHERAU	HYPOTHETICAL	68..4	KD P	5..13e+01
30	44	69..8	63..5	1	XNDPAEPO	ENDO-1,4-BETA-XYLANASE	5..13e+01		
31	44	69..8	109..1	1	NCA1CHICK	NEURAL CELL ADHESION M	5..13e+01		
32	43	68..3	103..0	1	SYB3MOUSE	SYNAPTOBREVIN 3 (CELLU	8..05e+01		
33	43	68..3	13..2	1	AYPEARATH	ATP SYNTHASE EPSILON C	8..05e+01		
34	43	68..3	21..6	1	FLA21METVA	FLAGELLIN B1 PRECURSOR	8..05e+01		
35	43	68..3	21..8	1	FLA1METVO	FLAGELLIN B1 PRECURSOR	8..05e+01		
36	43	68..3	22..0	1	Y069CAEEL	HYPOTHETICAL	2..9	KD P	8..05e+01
37	43	68..3	22..2	1	FLA21METVA	FLAGELLIN B2 PRECURSOR	8..05e+01		
38	43	68..3	34..2	1	AQPLHUMAN	AQUAPORIN-7 LIKE (AQUA	8..05e+01		
39	43	68..3	36..6	1	Y12L_SYNY3	HYPOTHETICAL	39..0	KD P	8..05e+01
40	43	68..3	39..4	1	FTSWHAEIN	CELL DIVISION PROTEIN	8..05e+01		
41	43	68..3	40..6	1	SGRAHPME	SERINE-GLYCOCYSTEYL AMI	8..05e+01		
42	43	68..3	49..3	1	ACHE_MOUSE	ACETYLCHOLINE RECEPTOR	8..05e+01		
43	43	68..3	53..6	1	FUJIFCAUR	M-RING PROTE	8..05e+01		
44	43	68..3	59..0	1	MDLA_ECOLI	MULTIDRUG RESISTANCE-L	8..05e+01		
45	43	68..3	110..8	1	GLANTYL_CYCLEASE	GC-E	8..05e+01		
					YAT				

ALIGNMENTS					
RESULT	1	HUMAN	STANDARD;	PRT;	118 AA.
ID	MAR1				
AC	Q16655;				
DT	01-NOV-1997	(Rel. 35, Created)			
DT	01-NOV-1997	(Rel. 35, Last sequence update)			
DT	15-JUL-1998	(Rel. 36, Last annotation update)			
DE	MELANOMA T-CELLS RECOGNIZED BY T-CELLS 1 (MART-1) (MELAN-A PROTEIN)				
DE	(ANTIGEN SK20-AA) (ANTIGEN LB39-AA).				
GN	MAR1				
OS	Homo sapiens (Human).				
OC	Bivalvia; Mollusca; Chordata; Craniata; Vertebrata; Mammalia;				
OC	Eutheria; Primates; Catarrhini; Hominidae; Homo.				
[1]					
RN	RP	SEQUENCE FROM N. A.			
RC	RC	TISSUE-MELANOMA;			
RX	RX	MEDLINE; 94224770.			
RA	RA	KAWAKAMI Y., ELIYAHU S., DELGADO C.H., ROBBINS P.F., RIVOLTINI L.,			
RA	RA	TOPALIAN S.L., MIKI T., ROSENBERG S.A.;			
RA	RA	"Cloning of the gene coding for a shared human melanoma antigen recognized by autologous T cells infiltrating into tumor.";			
RT	RT	Proc. Natl. Acad. Sci. U.S.A. 91:3515-3519(1994).			
RT	RT	[2]			
RT	RN	SEQUENCE FROM N. A.			
RP	RX	MEDLINE; 94275389.			
CC	CC	COULIE P.G., BRICHARD V., VAN PEL A., WOELFEL T., SCHNEIDER J.,			
CC	RA	TRAVERSARI C., MATTEI S., DE PLAEN E., LURQUIN C., SZIKORA J.-P.,			
CC	RA	RENAUD J.-C., BOON T.;			
CC	RA	"A new gene coding for a differentiation antigen recognized by autologous cytotoxic T lymphocytes on HLA-A2 melanomas."			
CC	RT	J. Exp. Med. 180:35-42(1994).			
CC	RL	-1- TISSUE SPECIFICITY: EXPRESSION IS RESTRICTED TO MELANOMA AND MELANOCTYE CELL LINES AND RETINA.			
CC	CC	-----			
CC	CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation of the European Bioinformatics Institute. There are no restrictions on use by non-profit institutions as long as its content is in no modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <a href="http://www.isb-sib.ch/announcements.html">http://www.isb-sib.ch/announcements.html</a> or send an email to licensed@isb-sib.ch).			
CC	CC	-----			
DR	DR	EMBL: U06455; AAA19238.1; -.			
DR	DR	EMBL: U06654; AAA20389.1; -.			
KW	KW	Antigen; Transmembrane.			
PT	PT	TRANSMEM			
		-----			
		POTENTIAL.			
		-----			
		27			



"Transcriptional analysis of the Streptococcus mutans hrca, grpe and dnaK genes and regulation of expression in response to heat shock and environmental acidification." Mol. Microbiol. 25:329-341(1997).

-1- FUNCTION: NEGATIVE REGULATOR OF CLASS I HEAT SHOCK GENES (GRPE-DNAK-DNAJ AND GROEL/GROES OPERONS). PREVENTS HEAT-SHOCK INDUCTION OF THESE OPERONS (BY SIMILARITY).

-1- SIMILARITY: BELONGS TO THE HRCA FAMILY.

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```

EMBL; U78296; AAC55610.1; -
PFAM; PF01628; HrcA; 1.
KW transcription regulation; Repressor; Heat shock.
SEQUENCE 345 AA; 39/06 MW; A0B065C2 CRC32;
Query Match 77.8%; Score 49; DB 1; Length 345;
Best Local Similarity 87.5%; pred. No. 4.74e+00;
Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

```

RESULT	5				
ID	YDGH_BACSU	STANDARD;	PRT;	885	AA.
AC	P96706;				
DT	15-DEC-1999	(Rel. 39, Created)			
DT	15-DEC-1999	(Rel. 39, Last sequence update)			
DT	15-DEC-1999	(Rel. 39, Last annotation update)			
DE	PUTATIVE MEMBRANE PROTEIN YDGH.				
GN	YDGH.				
DS	Bacillus subtilis.				
	Bacteria; Firmicutes; Bacillus/Clostridium group;				
DOC	Bacillus/Staphylococcus group; Bacillus				
	firmicutes; Firmicutes group				

SEQUENCE FROM N.A.  
STRAIN=168;  
KASAHIKA Y., NAKAI S., LEE S., SADAIE Y., OGASAWARA N.;  
A 148 kbp sequence of the region between 35 and 47 degree of the  
Bacillus genome;"  
Submitted (MAR-1987) to the EMBL/GenBank/DBJ databases.  
1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).  
-1- SIMILARITY: BELONGS TO THE MAMP FAMILY.

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or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch).

DR	EMBL; AB001488;	BAA1998..1;	-.
DR	EMBL; 299106;	CA12372..1;	-.
DR	EMBL; 299107;	CA12384..1;	-.
DR	SUBMITLIST;	BG12175;	YDGH.
KW	Hypothetical protein; Transmembrane.		
TRANSMEM	9	29	POTENTIAL.
TRANSMEM	181	201	POTENTIAL.
TRANSMEM	202	222	POTENTIAL.
TRANSMEM	227	247	POTENTIAL.
TRANSMEM	278	298	POTENTIAL.
TRANSMEM	304	324	POTENTIAL.
TRANSMEM	354	374	POTENTIAL.
TRANSMEM	716	736	POTENTIAL.

FT	TRANSNEM	740	760	POTENTIAL.
FT	TRANSNEM	772	792	POTENTIAL.
FT	TRANSNEM	817	837	POTENTIAL.
FT	TRANSNEM	847	867	POTENTIAL.
SQ	SEQUENCE	885 AA;	95488 MW;	4106171D CRC32;
Query Match	Score 49;	DB 1;	Length 885;	
Best Local Similarity	60.0%	Pred. No. 4.74e+00;		
Matches 6;	Conservative	2; Mismatches 2;	Indels 0;	Gaps 0;
Db	305 AVGGTLMII	314		
Qy	1 AAGIGLTIVI	10		
RESULT	6			
ID	YSA1_YEAST	STANDARD;	PRT;	231 AA.
AC	Q01976;			
DT	01-OCT-1993	(Rel. 27, Created)		
DT	01-OCT-1994	(Rel. 30, Last sequence update)		
DT	01-OCT-1996	(Rel. 34, Last annotation update)		
DE	YSA1 PROTEIN			
GN	YSA1 OR YBR111C OR YBR0907.			
OS	Eukaryotes cerevisiae (Baker's yeast).			
OC	Fungi: Ascomycota: Hemiascomycetes;			
OC	Saccharomycetaceae; Saccharomyces.			
RN	[1]			
RP	SEQUENCE FROM N.A.			
RC	SEQUIN= S288C;			
RX	MEDLINE: 932327848.			
RA	MANNHAUPT G., STUCKA R., EHNLE S., VETTER L., FELDMANN H.,			
RT	"Molecular analysis of yeast chromosome II between CMD1 and LYS2: the			
RT	excision repair gene RAD16 located in this region belongs to a novel			
RT	group of double-finger proteins.",			
RL	Yeast 8:397-408(1992).			
CC	-1 - SIMILARITY: STRONG. TO B SUBTILIS YQKG.			
CC	-1 - SIMILARITY: TO PROTEINS WITH A CORE MUTT DOMAIN.			
CC	-----			
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration			
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CC	-----			
DR	Z35980; CAA85068; 1; -.			
DR	X78993; CAA55614; 1; -.			
DR	X66247; CAA46972; 1; -.			
PIR	S44691; S44691.			
DR	SGD; L0002551; YSA1.			
DR	PROSITE; PS00893; MUTT; 1.			
DR	PF00293; mutt; 1.			
FT	DOMAIN 112 145. MUTT-LIKE.			
SQ	SEQUENCE 231 AA;	26087 MW;	49A2D6CB CRC32;	
Query Match	Score 48;	DB 1;	Length 231;	
Best Local Similarity	75.0%	Pred. No. 7.76e+00;		
Matches 6;	Conservative	2; Mismatches 0;	Indels 0;	Gaps 0;
Db	79 GIGILTIL 86			
Qy	1     ;			
Qy	3 GIGILTIL 10			
RESULT	7			

ID	Y4TP_RHISN	STANDARD;	PRT;	313 AA.	GN	RV0346C OR MTCY13E10.06C.
AC	Q53191;				OS	Mycobacterium tuberculosis.
DT	01-NOV-1997 (Rel. 35, Created)				OC	Bacteria; Firmicutes; Actinobacteria; Actinomycetidae;
DT	01-NOV-1997 (Rel. 35, Last annotation update)				OC	Corynebacterineae; Mycobacteriaceae; Mycobacterium.
DE	PROBABLE PEPTIDE ABC TRANSPORTER PERMEASE PROTEIN Y4TP.				RN	[1]
GN	OS	Rhizobium sp. (strain NGR234).			RP	SEQUENCE FROM N.A.
OG	Plasmid sym pNGR234a.				RC	SPRAIN-H37RV;
OC	Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;				RA	COLE S.T., BROSCHE R., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,
OC	Rhizobiaceae; Rhizobium.				RA	GORDON S.V., EIGENMEIR K., GAS C.E. III., BARRY C.E. F., TEKATA F.,
RN	[1]				RA	BADDOCK K., BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R.,
RP	SEQUENCE FROM N.A.				RA	DAVIES R., DEVLIN K., FELTMILL T., GENTLES S., HAMILIN N., HOLROYD S.,
RX	MEDLINE; 97305956.				RA	HORNSBY T., JAGELS K., KROGH A., MCLEAN J., MOULE S., MURPHY L.,
RA	FREIBERG C.A., FELLAY R., BAIROCH A., BROUGHTON W.J., ROSENTHAL A.,				RA	OLIVER S., OSBORNE J., RAJANDRAM M.A., ROGERS J.,
RA	PERET X.,				RA	RUTTER S., SEEGER K., SKELDON S., SQUIRES S., SOARES R., SULSTON J.E.,
RT	"Molecular basis of symbiosis between Rhizobium and legumes".				RA	TAYLOR K., WHITEHEAD S., BARRELL B.G.;
RL	Nature 387:394-401(1997).				RT	*Deciphering the biology of Mycobacterium tuberculosis from the complete genome sequence.";
RN	[2]				RL	Nature 393:537-544 (1998).
RP	SEQUENCE OF 107-313 FROM N.A.				CC	-1- FUNCTION: PROBABLE AMINO-ACID OR METABOLITE TRANSPORT PROTEIN.
RX	MEDLINE; 963889014.				CC	-1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
RA	FREIBERG C., PERET X., BROUGHTON W.J., ROSENTHAL A.;				CC	-1- SIMILARITY: BELONGS TO THE AMINO ACID PERMEASE FAMILY.
RT	"Sequencing the 500-kb GC-rich symbiotic replicon of Rhizobium sp. NGR234 using dye terminators and a thermostable 'sequenase': a beginning.";				CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <a href="http://www.isb-sib.ch/announce/">http://www.isb-sib.ch/announce/</a> or send an email to license@isb-sib.ch).
RT	Genome Res. 6:590-600(1996).				CC	DR EMBL: Z95324; CAB08578.1;
CC	-1- FUNCTION: PROBABLY PART OF A BINDING-PROTEIN-DEPENDENT TRANSPORT SYSTEM YATOPORS FOR A PEPTIDE. PROBABLY RESPONSIBLE FOR THE TRANSLOCATION OF THE SUBSTRATE ACROSS THE MEMBRANE.				DR PROSITE; PS00218; AMINO_ACID_PERMEASE; 1.	
CC	-1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. INNER MEMBRANE (POTENTIAL).				DR PFM: PF00324; aa_permeases; 1.	
CC	-1- SIMILARITY: WITH INTEGRAL MEMBRANE COMPONENTS OF OTHER BINDING-PROTEIN-DEPENDENT TRANSPORT SYSTEMS. BELONGS TO THE OPPBC SUBFAMILY.				DR PFM: PF00324; aa_permeases; 1.	
CC	This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <a href="http://www.isb-sib.ch/announce/">http://www.isb-sib.ch/announce/</a> or send an email to license@isb-sib.ch).				KW Hypothetical protein; Transport; Amino-acid transport; Transmembrane.	
CC	DR EMBL: AE000098; AAB91869.1;				FT TRANSMEM 26 46 POTENTIAL.	
DR	PROSITE; PS00402; BPD_TRANSP_INN_MEMBR; 1.				FT TRANSMEM 50 70 POTENTIAL.	
DR	PFAM; PF00528; BPD_transp; 1.				FT TRANSMEM 98 118 POTENTIAL.	
KW	Hypothetical protein; transport; Amino-acid transport; Transmembrane;				FT TRANSMEM 133 153 POTENTIAL.	
KW	Inner membrane; Plasmid.				FT TRANSMEM 163 183 POTENTIAL.	
FT	TRANSMEM 9 29 POTENTIAL.				FT TRANSMEM 214 234 POTENTIAL.	
FT	TRANSMEM 101 121 POTENTIAL.				FT TRANSMEM 256 276 POTENTIAL.	
FT	TRANSMEM 137 157 POTENTIAL.				FT TRANSMEM 290 310 POTENTIAL.	
FT	TRANSMEM 177 197 POTENTIAL.				FT TRANSMEM 341 361 POTENTIAL.	
FT	TRANSMEM 236 256 POTENTIAL.				FT TRANSMEM 369 389 POTENTIAL.	
FT	TRANSMEM 280 300 POTENTIAL.				FT TRANSMEM 414 434 POTENTIAL.	
SQ	SEQUENCE 313 AA; 34042 MW; 31F5F704 CRC32;				FT TRANSMEM 440 460 POTENTIAL.	
Query Match	Score 48; DB 1; Length 313;				SQ SEQUENCE 487 AA; 52194 MW; 64BBBBCC CRC32;	
Best Local Similarity	76.2%				Query Match 76.2%; Score 48; DB 1; Length 487;	
Matches	7; Conservative				Best Local Similarity 60.0%; Pred. No. 7.78e+00;	
Matches	1; Mismatches 1; Indels 0; Gaps 0;				Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;	
Db	235 AGVPILTVI 243				RESULT 9 ID Y561_HAEIN STANDARD; PRT; 633 AA.	
Qy	1 : : : : : : : : : :				AC P44016;	
Db	2 AGIGILTVI 10				DT 01-NOV-1995 (Rel. 32, Created)	
Qy					DT 01-NOV-1997 (Rel. 35, Last sequence update)	
Db	235 AGVPILTVI 243				DT 15-JUL-1998 (Rel. 36, Last annotation update)	
Qy	1 : : : : : : : : : :				DE HYPOTHETICAL PROTEIN HI0561/560.	
OS	Hemophilus influenzae; Proteobacteria; gamma subdivision; Pasteurellaceae;				GN HI0561/560	
OC	Bacteria; Hemophilus.				OS Hemophilus influenzae; Proteobacteria; gamma subdivision; Pasteurellaceae;	
OC	Hemophilus.				AC Hemophilus.	
RN	[1]				AC [1]	
RP	SEQUENCE FROM N.A.				DE SEQUENCE FROM N.A.	
RC	SPRAIN-FRD / KW20;				GN SPRAIN-FRD / KW20;	
RX	MEDLINE; 95350630.				OS MEDLINE; 95350630.	

RA FLEISCHMANN R.D., ADAMS M.D., WHITE O., CLAYTON R.A., KIRKNESS E.F.,  
 RA KERLAVAGE A.R., BULT C.J., TOMB J.-F., DOUGHERTY B.A., MERRICK J.M.,  
 RA MCKENNEY K., SUTTON G., FITZHUGH W., FIELDS C.A., GOCAYNE J.D.,  
 RA SCOTT J.D., SHIRLEY R., LIU L.-I., GLODEK A., KELLEY J.M.,  
 RA WEIDMAN J.F., PHILLIPS M.C.A., SPRIGGS T., HEDBLUM B., COFFON M.D.,  
 RA OTTERBACK T.R., HANNA M.C., NGUYEN D.T., SAUDEK D.M., BRANDON R.C.,  
 RA FINE L.D., FRITCHMAN J.L., FUHRMANN J.L., GEOGHAGEN N.S.M.,  
 RA GNEHM C.L., McDONALD L.A., SMALL K.V., FRASER C.M., SMITH H.O.,  
 RA VENTER J.C.;  
 RP REVISIONS.  
 RT "Whole-genome random sequencing and assembly of Haemophilus  
 influenzae Rd.";  
 RL Science 269:496-512(1995);  
 [2]

RA Submitted (SEP-1996) to the EMBL/GenBank/DDBJ databases.  
 RL -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).  
 CC

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 CC

DR EMBL; U32727; AC22215.1; -.

DR TIGR; HI0561; -.

RW HYPOTHETICAL protein; Transmembrane; POTENTIAL.

FT TRANSEM 8 28 POTENTIAL.

FT TRANSEM 45 65 POTENTIAL.

FT TRANSEM 70 90 POTENTIAL.

FT TRANSEM 128 148 POTENTIAL.

FT TRANSEM 180 200 POTENTIAL.

FT TRANSEM 230 250 POTENTIAL.

FT TRANSEM 281 301 POTENTIAL.

FT TRANSEM 311 331 POTENTIAL.

FT TRANSEM 345 365 POTENTIAL.

FT TRANSEM 379 399 POTENTIAL.

FT TRANSEM 420 440 POTENTIAL.

FT TRANSEM 483 503 POTENTIAL.

FT TRANSEM 515 535 POTENTIAL.

FT TRANSEM 564 584 POTENTIAL.

FT TRANSEM 604 624 POTENTIAL.

SQ SEQUENCE 633 AA; 6616 MW; A940DDD CRC32;

Query Match Score 48; DB 1; Length 633;  
 Best Local Similarity 66.7%; Pred. No. 7.76e+00;  
 Matches 6; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Db 347 SGIGLSVI 355  
 QY :||||:||| 2 AGIGLTVI 10

RESULT 10  
 ID CYAB\_1EIDO STANDARD; PRT; 1331 AA.  
 AC Q25264;  
 DT 15-JUL-1998 (Rel. 36, Created)  
 DT 15-JUL-1998 (Rel. 36, Last sequence update)  
 DT 15-DEC-1998 (Rel. 37, Last annotation update)  
 DE RECEPTOR-TYPE ADENYLATE CYCLASE B (EC 4.6.1.1) (ATP PYROPHOSPHATE-  
 DE LIASE (ADENYLIC CYCLASE), RAC-B)  
 GN Leishmania donovani.  
 CS Leishmania donovani; Kinetoplastida; Trypanosomatidae; Leishmania.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-15 SUDANSE;  
 RX MEDLINE; 9534054.  
 RA SANCHEZ M.A., ZEOLI D., KIAMO E.M., KAVANAUGH M.P., LANDFEAR S.M.;  
 RT "A family of putative receptor-adenylate cyclases from Leishmania  
 donovani.";  
 RT

RL J. Biol. Chem. 270:17551-17558(1995).  
 CC COULD ACT AS A RECEPTOR FOR A UNKNOWN LIGAND.  
 CC -1- CATALYTIC ACTIVITY: ATP = 3',5'-CYCLOC AMP + PYROPHOSPHATE.  
 CC -1- SUBCELLULAR LOCATION: MEMBRANE-BOUND.  
 CC -1- DEVELOPMENTAL STAGE: EXPRESSED IN THE INSECT STAGE (PROMASTIGOTE)  
 CC BUT NOT IN THE MAMMALIAN HOST STAGE OF THE PARASITE LIFE CYCLE.  
 CC -1- SIMILARITY: BELONGS TO ADENYLIC CYCLASE CLASS 3 FAMILY.  
 CC  
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 CC or send an email to license@isb-sib.ch).  
 CC  
 DR PFAM; PF00211; quanyalate\_cyc\_1.  
 DR PFM: PF00211; quanyalate\_cyc\_1.  
 KW Lyase; CAMP synthetis; Transmembrane; Receptor; Glycoprotein.  
 FT DOMAIN 1 33 CYTOPLASMIC (POTENTIAL).  
 FT DOMAIN 34 54 POTENTIAL.  
 FT DOMAIN 55 898 EXTRACELLULAR (POTENTIAL).  
 FT DOMAIN 899 919 POTENTIAL.  
 FT DOMAIN 920 1331 CYTOPLASMIC (POTENTIAL).  
 FT CARBOHYD 255 255 POTENTIAL.  
 FT CARBOHYD 429 429 POTENTIAL.  
 FT CARBOHYD 558 558 POTENTIAL.  
 FT CARBOHYD 574 574 POTENTIAL.  
 FT CARBOHYD 657 657 POTENTIAL.  
 SQ SEQUENCE 1331 AA; 144162 MW; CCC01FC9 CRC32;  
 DR 902 AGIALLTVI 910  
 DR ||||:|||||  
 DR 2 AGIGLTVI 10  
 DR 902 AGIALLTVI 910  
 DR ||||:|||||  
 DR 2 AGIGLTVI 10  
 DR RESULT 11  
 ID ATPL\_SULAC STANDARD; PRT; 101 AA.  
 AC P23040;  
 DT 01-NOV-1991 (Rel. 20, Created)  
 DT 01-NOV-1991 (Rel. 20, Last sequence update)  
 DT 15-JUL-1998 (Rel. 36, Last annotation update)  
 DE MEMBRANE-ASSOCIATED ATPASE C CHAIN (EC 3.6.1.34) (SUL-ATPASE-  
 DE PROTEOLIPID CHAIN).  
 GN ATPP.  
 OS Sulfolobus acidocaldarius.  
 OC RN [1]  
 RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.  
 RX MEDLINE; 89214142.  
 RA DENDA K., KONISHI J., OSHIMA T., DATE T., YOSHIDA M.;  
 RA "A gene encoding the proteolipid subunit of Sulfolobus acidocaldarius  
 RT ATPase complex";  
 RL J. Biol. Chem. 264:7119-7121(1989).  
 CC -1- FUNCTION: THE C CHAIN IS A PROTEOLIPID, AND ONE OF THE MEMBRANOUS  
 CC SUBUNITS OF THE THE NONENZYMATIC COMPONENT OF THE SUL-ATPASE  
 CC COMPLEX.  
 CC -1- SUBUNIT: SUL-ATPASE IS COMPOSED OF SIX (OR FIVE ?) SUBUNITS;  
 CC ALPHA, BETA, GAMMA, C (PROTEOLIPID), AND POSSIBLY EPSILON.  
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).  
 CC -1- SIMILARITY: BELONGS TO THE V-ATPASE PROTEOLIPID SUBUNIT FAMILY.  
 CC  
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 CC or send an email to license@isb-sib.ch).

CC EMBL; J04740; AAAT72/03.1; -  
 DR PIR; A33351; A33351.  
 DR HSSP; P00138; ICGN.  
 DR PFAM; PF00137; ATP-synt\_C; 1.  
 KW Hydrogen ion transport; Lipid-binding; Transmembrane.  
 FT TRANSMEM 5 25 POTENTIAL.  
 FT TRANSMEM 37 57 POTENTIAL.  
 FT TRANSMEM 75 95 POTENTIAL.  
 SQ SEQUENCE 101 AA; 10362 MW; 1DC8<74D CRC32;

Query Match Score 47; DB 1; Length 308;  
 Best Local Similarity 87.5%; Pred. No. 1.26e+01;  
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Db 59 AAGIGVLT 66  
 QY 1 AAGIGVLT 8

RESULT 12 MENA\_HAEIN STANDARD; PRT; 308 AA.  
 ID P44759; 01-NOV-1995 (Rel. 32, Created)  
 DT 01-NOV-1995 (Rel. 32, Last sequence update)  
 DT 15-JUL-1998 (Rel. 36, Last annotation update)  
 DE 1,4-DIHYDROXY-2-NAPHTHOATE OCTAPRENYLTRANSFERASE (EC 2.5.1.-) (DHNA-OCTAPRENYLTRANSFERASE).  
 GN OS Haemophilus influenzae.  
 OS Haemophilus influenzae.  
 OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX STRAIN=RD / KWA.  
 RA FLEISCHMANN R.D., ADAMS M.D., WHITE O., CLAYTON R.A., KIRKNESS E.F., SUTTON G.G., STRAIN=RD / KWA.  
 RA MEDLINE; 95350650.  
 RA KERLAVERAGE A.R., BULT C.J., TOMB J.-P., DOUGHERTY B.A., MERRICK J.M., MCKENNEY K., SUTTON G., FITZGERALD L.M., LEWIS N., ADAMS M.D., HICKEY E.K., MCNEILAN K., QUACKENBUSH J.J., ZHOU L., KIRKNESS E.F., DOUGHERTY B.A., LOFTUS B., RICHARDSON D., DODSON R., KHALAK H.G., GLODEK A., MCKENNEY K., FITZGERALD L.M., LEWIS N., ADAMS M.D., HICKEY E.K., BERG D.E., GOCayne J.D., UTTERBACK T.R., PETERSON J.D., KELLEY J.M., COPTON M.D., WEIDMAN J.M., FUJI T.C., BOWMAN C., WATTHAY L., WALLIN E., HAYES W.S., BORODovsky M., KARP P.D., SMITH H.O., FRASER C.M., VENTER J.C.; "The complete genome sequence of the gastric pathogen Helicobacter pylori"; RT Nature 388:539-547 (1997).  
 RA "Whole-genome random sequencing and assembly of Haemophilus influenzae Rd"; RT "Whole-genome random sequencing and assembly of Haemophilus influenzae Rd"; RT Science 265:496-512 (1995).  
 CC -I- FUNCTION: CONVERSION OF 1,4-DIHYDROXY-2-NAPHTHOATE (DHNA) TO DIMETHYLMENAQUNONE (DMQ). ATTACHES OCTAPRENYLYLPOPHOSPHATE, A MEMBRANE-BOUND 40-CARBON SIDE CHAIN TO DHNA. THE CONVERSION OF DHNA TO DMQ PROCEEDS IN THREE STAGES: THE REMOVAL OF THE CARBOXYL GROUP OF DHNA AS CO2, THE ATTACHMENT OF THE ISOPRENOID SIDE CHAIN, AND A QUINOI-TO-QUINONE OXIDATION, WHICH IS THOUGHT TO BE SPONTANEOUS (BY SIMILARITY).  
 CC -I- PATHWAY: MENAQUNONE BIOSYNTHESIS.  
 CC -I- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. INNER MEMBRANE (BY SIMILARITY).  
 CC -I- SIMILARITY: BELONGS TO THE MENA FAMILY.

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CC EMBL; U32732; AAC22167.1; -  
 DR TIGR; HI0509; -  
 KW Menaqunone biosynthesis; transferase; Transmembrane; Membrane.  
 FT TRANSMEM 22 42 POTENTIAL.  
 FT TRANSMEM 47 67 POTENTIAL.

FT TRANSMEM 101 121 POTENTIAL.  
 FT TRANSMEM 129 149 POTENTIAL.  
 FT TRANSMEM 153 173 POTENTIAL.  
 FT TRANSMEM 186 206 POTENTIAL.  
 FT TRANSMEM 235 255 POTENTIAL.  
 FT TRANSMEM 286 306 POTENTIAL.  
 SQ SEQUENCE 308 AA; 33345 MW; 090B2655 CRC32;

Query Match Score 47; DB 1; Length 308;  
 Best Local Similarity 55.6%; Pred. No. 1.26e+01;  
 Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Db 132 AGIGILAVI 140  
 QY 2 AGIGILAVI 10

RESULT 13 SECD\_HELPPY STANDARD; PRT; 503 AA.  
 ID SECD\_HELPPY  
 AC 026074;  
 DT 15-DEC-1999 (Rel. 39, Created)  
 DT 15-DEC-1999 (Rel. 39, Last sequence update)  
 DE PRONEIN-EXPORT MEMBRANE PROTEIN SECD.  
 GN SECD OR HP1550.  
 OS Helicobacter pylori (Campylobacter pylori); Bacteria; Proteobacteria; epsilon subdivision; Helicobacter group; Helicobacter.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX STRAIN=66695 / ATCC 700392;  
 RA MEDLINE; 97394467.  
 RA TOMB J.-F., WHITE O., KERLAVERAGE A.R., CLAYTON R.A., SUTTON G.G., FLEISCHMANN R.D., KETCHUM K.A., KLENK H.-P., GILL S., DOUGHERTY B.A., NELSON K., QUACKENBUSH J.J., ZHOU L., KIRKNESS E.F., PETERSON S., RA MEDLINE; 97394467.  
 RA LOFTUS B., RICHARDSON D., DODSON R., KHALAK H.G., GLODEK A., RA MCKENNEY K., FITZGERALD L.M., LEWIS N., ADAMS M.D., HICKEY E.K., RA BERG D.E., GOCayne J.D., UTTERBACK T.R., PETERSON J.D., KELLEY J.M., RA COPTON M.D., WEIDMAN J.M., FUJI T.C., BOWMAN C., WATTHAY L., WALLIN E., RA HAYES W.S., BORODovsky M., KARP P.D., SMITH H.O., FRASER C.M., RA VENTER J.C.; "The complete genome sequence of the gastric pathogen Helicobacter pylori"; RT Nature 388:539-547 (1997).  
 CC -I- FUNCTION: INVOLVED IN PROTEIN EXPORT (BY SIMILARITY).  
 CC -I- SUBUNIT: PART OF THE PROKARYOTIC PROTEIN TRANSLOCATION APPARATUS WHICH COMPRISE SECA, SECB, SECY, SECZ, SECG AND SECY (BY SIMILARITY).  
 CC -I- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).  
 CC -I- SIMILARITY: BELONGS TO THE SECD/SECY FAMILY. SECY FAMILY.  
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 CC EMBL; AB000652; AAD08588.1; -  
 DR TIGR; HP1550; -  
 KW Protein transport; Translocation; Transmembrane; Membrane.  
 FT TRANSMEM 199 219 POTENTIAL.  
 FT TRANSMEM 334 354 POTENTIAL.  
 FT TRANSMEM 357 377 POTENTIAL.  
 FT TRANSMEM 383 403 POTENTIAL.  
 FT TRANSMEM 456 476 POTENTIAL.  
 SQ SEQUENCE 503 AA; 54247 MW; 9A76592C CRC32;

Query Match Score 47; DB 1; Length 503;  
 Best Local Similarity 60.0%; Pred. No. 1.26e+01;  
 Matches 3; Mismatches 1; Indels 0; Gaps 0;

Db 460 TTGIGILASI 469  
Qy :|||||: | 1 AAGIGILTVI 10

RESULT 14  
ID SECD\_HELPJ STANDARD; PRT; 526 AA.  
AC Q9ZL66;  
DT 15-DEC-1999 (Rel. 39, Created)  
DT 15-DEC-1999 (Rel. 39, Last sequence update)  
DT 15-DEC-1999 (Rel. 39, Last annotation update)  
DE PROTEIN-EXPORT MEMBRANE PROTEIN.  
GN SECD.  
OS Helicobacter pylori J99 (campylobacter pylori J99).  
OC Bacteria: Proteobacteria; epsilon subdivision; Helicobacter group;  
OC Helicobacter.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE; 99120557.  
RA ALM R.A., LING L.-S.L., MOIR D.T., KING B.L., BROWN E.D., DOIG P.C.,  
RA SMITH D.R., NOONAN B., GUILDFORD B.C., DEJONCE B.L., CARMEL G.,  
RA TUMMINO P.J., CARUSO A., URIA-NICKELSEN M., MILLS D.M., IVES C.,  
RA TRUST T.J.;  
RT "Genomic sequence comparison of two unrelated isolates of the human  
gastric pathogen Helicobacter pylori.";  
RL Nature 397:1786-1780(1999).  
CC -1- FUNCTION: INVOLVED IN PROTEIN EXPORT (BY SIMILARITY).  
CC -1- SUBUNIT: PART OF THE PROKARYOTIC PROTEIN TRANSPORT APPARATUS  
CC WHICH COMPRISE SECA, SECB, SECY, SECCE, SECF, SECG AND SECY  
CC (BY SIMILARITY).  
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (BY SIMILARITY).  
CC -1- SIMILARITY: BELONGS TO THE SECDF/SECY FAMILY. SECDF FAMILY.  
CC DR EMBL; AE001567; AAD07024\_1; -  
CC KW Protein transport; Translocation; Membrane.  
CC FT TRANSMEM 8 28 POTENTIAL.  
CC FT TRANSMEM 356 376 POTENTIAL.  
CC FT TRANSMEM 379 399 POTENTIAL.  
CC FT TRANSMEM 453 473 POTENTIAL.  
CC FT TRANSMEM 478 498 POTENTIAL.  
CC SEQUENCE 526 AA; 56796 MW; FCDEBF9A9 CRC32;

RX MEDLINE; 98344137.  
RA KAWARABAYASI Y., SAWADA M., HORIKAWA H., HAKIWA Y., HINO Y.,  
RA YAMAMOTO S., SEKINE M., BABA S., KOSUGI H., HOSOYAMA A., NAGAI Y.,  
RA SAKAI M., OGURA K., OTSUKA R., NAKAZAWA H., TAKAMIYA M., OHFUKU Y.,  
RA FUNAHASHI T., TANAKA T., KUDOH Y., YAMAZAKI J., KUSHIDA N., OGUCHI A.,  
RA AOKI K., NAKAMURA Y., HORIKOSHI K., MASUCHI Y., SHIZUYA H.,  
RA KIKUCHI H.;  
RT "Complete sequence and gene organization of the genome of a hyper-  
thermophilic archaeabacterium, Pyrococcus horikoshii, OT3.";  
RT DNA Res. 5:55-76(1998).  
RL -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).  
CC -1 - SIMILARITY: BELONGS TO THE UDP0056 (MARC) FAMILY.  
CC -1-  
CC This SWISS-PROT entry is copyright. It is produced through a collaboration  
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -  
CC the European Bioinformatics Institute. There are no restrictions on its  
CC use by non-profit institutions as long as its content is in no way  
CC modified and this statement is not removed. Usage by and for commercial  
CC entities requires a license agreement. (See http://www.isb-sib.ch/announce/  
CC -1-  
CC EMBL; AP00003; BAA29851.1; -  
CC DR EMBL; AP00003; BAA29851.1; -  
CC KW Hypothetical protein; Transmembrane.  
FT TRANSMEM 8 28 POTENTIAL.  
FT TRANSMEM 49 69 POTENTIAL.  
FT TRANSMEM 73 93 POTENTIAL.  
FT TRANSMEM 111 131 POTENTIAL.  
FT TRANSMEM 140 160 POTENTIAL.  
FT TRANSMEM 181 201 AA; 21592 MW; 97675186 CRC32;  
SQ SEQUENCE 201 AA; 21592 MW; 97675186 CRC32;

Query Match Score 46; DB 1; Length 201;  
Best Local Similarity 60.0%; Pred. No. 2.03e-01;  
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Db 138 SPGIVILVII 147  
Qy ::||| |||: | 1 AAGIGILTVI 10

Search completed: Fri May 5 22:18:20 2000  
Job time : 39 secs.

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Db 482 TTGIGILASI 491  
Qy :|||||: | 1 AAGIGILTVI 10

RESULT 15  
ID Y760\_PYRHO STANDARD; PRT; 201 AA.  
AC 058499;  
DT 15-DEC-1998 (Rel. 37, Created)  
DT 15-DEC-1998 (Rel. 37, Last sequence update)  
DT 15-DEC-1999 (Rel. 39, Last annotation update)  
DE HYPOTHETICAL PROTEIN PH0760.  
GN PH0760 OR PHC1026.  
OS Pyrococcus horikoshii.  
OC Archaea; Euryarchaeota; Thermococcales; Thermococcaceae; Pyrococcus.  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=OT3;



QY 1 AAGIGILTVI 10  
 RESULT 2 VGBEPB #type complete  
 ENTRY glycoprotein gIII precursor - suid herpesvirus 1  
 TITLE formal\_name suid herpesvirus 1  
 ORGANISM 30-Sep-1987 #sequence\_revision 30-Sep-1987 #text\_change  
 DATE 16-Jul-1999  
 ACCESSION A26097  
 REFERENCE Robbins, A.K.; Watson, R.J.; Whealy, M.E.; Hays, W.W.; Enquist, L.W.  
 #authors J. Virol. (1986) 58:339-347  
 #journal Characterization of a Pseudorabies virus glycoprotein gene with homology to herpes simplex virus type 1 and type 2 glycoprotein C.  
 #cross-references MUID:86200375  
 #accession A26097  
 #molecule\_type DNA  
 ##residues 1-479 ##label ROB  
 #cross-references GB:MI2778; NID:9334049; PID:9334050  
 #experimental\_source strain Becker  
 CLASSIFICATION #superfamily herpesvirus glycoprotein F  
 KEYWORDS glycoprotein  
 FEATURE 1-22 #domain signal sequence #status predicted #label SIG\\  
 23-479 #product glycoprotein gIII #status predicted #label GPG\\  
 40,84,169,192,220, #binding\_site carbohydrate (Asn) (covalent) #status  
 228,285,302 predicted  
 SUMMARY #length 479 #molecular\_weight 51206 #checksum 1630  
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 Matches 6; Conservative 3; Mismatches 0; Indels 0; Gaps 0;  
 Db 456 AGIGILAVI 64  
 QY 2 AGIGILTVI 10  
 RESULT 3 D70073 #type complete  
 ENTRY metabolite transport protein homolog yxcc - Bacillus subtilis  
 TITLE formal\_name Bacillus subtilis  
 ORGANISM 05-Dec-1997 #sequence\_revision 05-Dec-1997 #text\_change  
 DATE 24-Sep-1999  
 ACCESSION A69580  
 #authors Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Bertero, M.G.; Bessieres, P.; Bolotin, A.; Borchart, S.; Boris, R.; Bourrel, L.; Brans, A.; Braun, M.; Brignell, S.C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Choi, S.K.; Codani, J.J.; Connerton, I.F.; Cummings, N.J.; Daniel, R.A.; Denizot, F.; Devine, K.M.; Duesterhoeft, A.; Ehrlich, S.D.; Emmerson, P.T.; Entian, K.D.; Errington, J.; Fabret, C.; Ferrari, E.; Poulter, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Gallizzi, A.; Galloren, N.; Ghim, S.Y.; Glaser, P.; Goffeau, A.; Golightly, E.J.; Grandi, G.; Giuseppi, G.; Guy, B.J.; Haga, K.; Haiech, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Holzapfel, S.; Hosono, S.; Hull, M.F.; Itaya, M.; Jones, L.; Joris, B.; Karanam, D.; Kasahara, Y.; Klaerr-Blanchard, M.; Klein, C.; Kobayashi, Y.; Koetter, P.; Konigstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois, S.; Lauber, J.; LaJarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Mauel, C.; Medigue, C.; Medina, N.; Mellado, R.P.; Mizuno, M.; Moestl, D.; Nakai, S.; Noback, M.; Neone, D.; O'Reilly, N.; Ogawa, K.; Ogiwara, A.; Oudega, B.; Park, S.H.; Parro, V.; Pohl, T.M.; Portebeille, D.; Porvolik, S.; Prescott, A.M.; Presecan, E.; Pujic, P.; Purnelle, B.; Rapoport, G.; Rey, M.; Reynolds, S.; Rieger, M.; Rivolta, C.; Rocha, E.; Roche, B.; Rose, M.; Sadaie, Y.; Sato, T.; Scanlon, E.; Schleich, S.; Schroeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Seror, S.J.; Seror, P.; Shin, B.S.; Soldo, B.; Sorokin, A.; Tacconi, E.; Takagi, T.; Takahashi, H.; Takemaru, K.; Takeuchi, M.; Tamakoshi, A.; Tanaka, T.; Terpstra, P.; Tognoni, A.; Tosato, V.; Uchiyama, S.; Vandenberg, M.; Vanier, F.; Vassarotti, A.; Vilar, A.; Wanbutt, R.; Wedler, E.; Weitzneger, T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yata, K.; Yoshida, K.; Yoshihikawa, H.F.; Zumstein, E.; Yoshikawa, H.; Danchin, A.

#journal Nature (1997) 390:249-256  
 #title The complete genome sequence of the Gram-positive bacterium *Bacillus subtilis*.  
 #cross-references MUID:98044033  
 #accession D70073  
 #status preliminary; nucleic acid sequence not shown; translation not shown  
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 ##residues 1-461 ##label RUN  
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 #experimental\_source strain 168  
 GENETICS  
 #gene YXCC  
 CLASSIFICATION #superfamily glucose transport protein SUMMARY #length 461 #molecular\_weight 50140 #checksum 8642  
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 Best Local Similarity 87.5%; Pred. No. 9.80e+00;  
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 Db 285 GIGILAVI 292  
 Qy 3 GIGILTVI 10  
 RESULT 4 H70597 #type complete  
 ENTRY probable membrane protein - Mycobacterium tuberculosis  
 TITLE (strain H37RV)  
 ORGANISM #formal\_name Mycobacterium tuberculosis  
 DATE 17-Jul-1998 #sequence\_revision 17-Jul-1998  
 ACCESSION H70597  
 REFERENCE Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.V.; Eiglmeier, K.; Gas, S.; Barry III, C.E.; Tekwani, F.; Badcock, K.; Basham, D.; Brown, D.; Chillingworth, T.; Connor, R.; Davies, R.; Devin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Hornsby, T.; Jagels, K.; Krogh, A.; McLean, J.; Moule, S.; Murphy, L.; Oliver, S.; Osborne, J.; Quail, M.A.; Rajandream, M.A.; Rogers, J.; Rutte, S.; Seeger, K.; Skelton, S.; Squares, S.; Saarens, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.  
 #journal Nature (1998) 393:537-544  
 #title Deciphering the biology of Mycobacterium tuberculosis from the complete genome sequence.  
 #cross-references MUID:98295987  
 #accession H70597  
 #status preliminary; nucleic acid sequence not shown; translation not shown  
 #molecule\_type DNA  
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 PID:g1944601  
 #experimental\_source strain H37Rv  
 GENETICS  
 #gene RV3887C  
 SUMMARY #length 509 #molecular\_weight 53278 #checksum 6762

Query Match 79.4%; Score 50; DB 2; Length 509;  
Best Local Similarity 70.0%; Pred. No. 9.80e+00;  
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Db 172 AGGIGILVVI 181  
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Qy 1 AAGGILVVI 10

RESULT 5 T00502 #type complete  
Protein kinase homolog T20D16.7 - Arabidopsis thaliana  
#formal\_name Arabidopsis thaliana #common\_name mouse-ear  
cress  
01-Feb-1999 #sequence\_revision 01-Feb-1999 #text\_change  
T20D16.7  
Rounsey, S.D.; Lin, X.; Ketchum, K.A.; Crosby, M.L.;  
Brandon, R.C.; Sykes, S.M.; Kaul, S.; Mason, T.M.;  
Kharavage, A.R.; Adams, M.D.; Somerville, C.R.; Venter,  
J.C.  
submitted to the EMBL Data Library, November 1997  
Arabidopsis thaliana chromosome II BAC T20D16 genomic  
sequence.  
T00502  
##status translated from GB/EMBL/DDJB  
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##residues 1-773 #label R01  
##cross-references EMBL:AC02391; NID:92642427; PID:92642433  
##experimental\_source cultivar Columbia  
##map-position 2  
##introns 545/1  
##note T20D16.7  
##length 773 #molecular-weight 84148 #checksum 123

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Best Local Similarity 77.8%; Pred. No. 9.80e+00;  
Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Db 343 AGGIGIL 351  
|:|||:|||:  
Qy 2 AGGIGILVVI 10

RESULT 6 T15544 #type complete  
hypothetical protein C18A3.2 - Caenorhabditis elegans  
#formal\_name Caenorhabditis elegans #common\_name  
C. elegans  
20-Sep-1999 #sequence\_revision 20-Sep-1999 #text\_change  
T15544  
218367  
#accessions  
#authors Hallsworth, K.  
#submission submitted to the EMBL Data Library, June 1995  
#description The sequence of C. elegans cosmid C18A3.  
T15544  
#accession  
#status preliminary; translated from GB/EMBL/DDJB  
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PIDN:AA68371.1; CESP:C18A3.2  
##experimental\_source strain Bristol N2  
##genetics

RESULT 7 B69783 #type complete  
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#formal\_name Bacillus subtilis  
05-Dec-1997 #sequence\_revision 05-Dec-1997 #text\_change  
24-Sep-1998  
B69783  
A65580  
Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.;  
Alloni, G.; Azevedo, V.; Bertero, M.G.; Bessieres, P.;  
Bolotin, A.; Borchart, S.; Bories, R.; Boursier, L.; Branciforte, S.;  
A.; Braun, M.; Brignell, S.C.; Bron, S.; Brouillet, S.;  
Bruschi, C.V.; Caldwell, V.; Carter, N.M.;  
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Glaser, A.; Goffeau, A.; Golightly, E.J.; Grandi, G.;  
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M.; Ogawa, K.; Ogiwara, A.; Oudega, B.; Park, S.H.; Parro,  
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A.M.; Presecan, E.; Pujic, P.; Purnelle, B.; Rapoport, G.;  
Rey, M.; Reynolds, S.; Rieger, M.; Rivolta, C.; Rocha, E.;  
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Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto,  
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Yoshikawa, H.; Danchin, A.

#journal Nature (1997) 390:249-256  
#title The complete genome sequence of the Gram-positive bacterium  
#cross-references MURID:98044033  
#accession B69783  
##status preliminary; nucleic acid sequence not shown;  
translaton not shown

##molecule-type DNA  
##residues 1-885 ##label KUN  
##cross-references GB:Z99106; PID:e118254; PID:92632865;  
PID:e1182531; PID:92632865

GENETICS  
#experimental\_source strain 168  
#gene Ydgh  
SUMMARY  
#length 885 #molecular-weight 95488 #checksum 3557

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Best Local Similarity 70.0%; Pred. No. 1.54e+01;  
Matches 6; Conservative 2; Mismatches 0; Gaps 0;

Db 305 AVGYGLMII 314  
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Qy 1 AAGGILVVI 10

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RESULT          8          S48276  #type complete
ENTRY          TITL          YSA1 protein - yeast (Saccharomyces cerevisiae)
AUTENNATE_NAMES          protein YBR007; protein YBR11C
ORGANISM          #formal_name Saccharomyces cerevisiae
DATE          10-Sep-1999 *sequence_revision 10-Sep-1999 #text_change
10-Sep-1999
ACCESSIONS          S48276; S45979; S25364; S44691
REFERENCE          S48255
#authors          Maenhaut, G.; Stucka, R.; Ehnle, S.; Vetter, I.; Feldmann, H.; Yeast (1994) 10:1363-1381
#journal          Yeast (1994) 10:1363-1381
#title           Analysis of a 70 kb region on the right arm of yeast chromosome II.
#cross-references MUID:95208357
#accession        S48276
##status          nucleic acid sequence not shown
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REFERENCE          S45927
#authors          Feldmann, H.; Mannhaupt, G.; Schwarzlose, C.; Vetter, I.
#submission        submitted to the Protein Sequence Database, August 1994
#accession        S45979
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#cross-references EMBL:Z35980; NID:9536465; PID:9536466; MIPS:YBR11C
#accession        S25364
#authors          Maenhaut, G.; Stucka, R.; Ehnle, S.; Vetter, I.; Feldmann, H.; Yeast (1992) 8:397-408
#journal          Yeast (1992) 8:397-408
#title           Molecular analysis of yeast chromosome II between CMD1 and LYS2: the excision repair gene RAD16 located in this region belongs to a novel group of double-finger proteins.
#cross-references SGD:YSA1
#accession        S25364
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##residues         1-47 ##label MAW
#cross-references EMBL:X66247; NID:93548; PID:g3549
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#cross-references SGD:SC000315; MIPS:YBR11C
#map-position 2R
CLASSIFICATION          #superfamily YffH protein; mutT domain homology
FEATURE          111-145      #domain mutT domain homology #label MUTT
SUMMARY          231      #length 231      #molecular_weight 26687 #checksum 4809
Query Match          Best Local Similarity 76.2%; Score 48; DB 1; Length 231;
Matches          6; conservative 2; Mismatches 0; Indels 0; Gaps 0; RESULT          10
Db               79 GIGILTIL 86          C70574  #type complete
Qy               |||||:          probable aroP2 protein - Mycobacterium tuberculosis (strain H37Rv)
          3 GIGILTVI 10          C70574  #formal_name Mycobacterium tuberculosis (strain H37Rv)
          #text_change
RESULT          9          F69798  #type complete
ENTRY          TITL          conserved hypothetical protein yetI - Bacillus subtilis
AUTENNATE_NAMES          #formal_name Bacillus subtilis
DATE          05-Dec-1997 *sequence_revision 05-Dec-1997 #text_change
24-Sep-1998
ACCESSIONS          F69798
REFERENCE          A6980
#authors          Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Bertero, M.G.; Bessieres, P.; Blotoni, A.; Borchart, S.; Borsig, R.; Brouillet, S.; Brans, A.; Braun, M.; Brignell, S.C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Choi, S.K.; Codani, J.J.; Connerton, I.F.; Cummings, N.J.; Daniel, R.A.; Denizot, F.; Devine, K.M.; Duesterhoeft, A.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.V.; Eigemeier, K.; Gas, S.; Barry III, C.E.; Tekla, F.; Badcock, K.; Basham, D.; Brown, D.; Chillingworth, T.; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Hornby, L.; Oliver, S.; Osborne, J.; Quail, M.A.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squars, S.; Soares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
#journal          Nature (1998) 393:537-544
#text_change

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#title Deciphering the biology of *Mycobacterium tuberculosis* from  
the complete genome sequence.  
#cross-references MUID:98295987

#accession C70574 #status preliminary; nucleic acid sequence not shown;  
translation not shown

##molecule\_type DNA  
##residues 1-487 ##label COL  
##cross-references GB:295324; GB:AL123456; NID:93261760;  
PID:e315461; PID:92094825  
##experimental\_source strain H37Rv

GENETICS  
#gene arop2  
CLASSIFICATION #superfamily arginine permease  
SUMMARY #length 487 #molecular-weight 52194 #checksum 4052

Query Match 76.2%; Score 48; DB 2; Length 487;  
Best Local Similarity 60.0%; Pred. No. 2.40e+01;  
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Db 351 TAGIGLIGI 360  
QY 1 AAGIGILTVI 10

RESULT 11  
ENTRY F70682 #type complete  
probable membrane protein - *Mycobacterium tuberculosis*  
(strain H37RV)  
ORGANISM #formal\_name *Mycobacterium tuberculosis*  
DATE 17-Jul-1998 #sequence\_revision 17-Jul-1998 #text\_change  
17-Jul-1998  
ACCESSIONS F70682  
REFERENCE Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.V.; Eiglmeier, K.; Gas, S.; Barry, III, C.E.; Tekala, F.; Badcock, K.; Basham, D.; Brown, D.; Chillingworth, T.; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Hornsby, T.; Jagels, K.; Krogh, A.; McLean, J.; Moule, S.; Murphy, L.; Oliver, S.; Osborne, J.; Quail, M.A.; Rajandream, M.A.; Rogers, J.; Rutte, S.; Seeger, K.; Skelton, S.; Squares, S.; squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.  
#journal Nature (1998) 393:537-544  
#title Deciphering the biology of *Mycobacterium tuberculosis* from  
the complete genome sequence.  
#cross-references MUID:98295987

#accession F70682 #status preliminary; nucleic acid sequence not shown;  
translation not shown

##molecule\_type DNA  
##residues 1-667 ##label COL  
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##experimental\_source strain H37Rv

GENETICS  
#gene Rv2395  
SUMMARY #length 667 #molecular-weight 68251 #checksum 9102

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Best Local Similarity 55.6%; Pred. No. 2.40e+01;  
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Df 379 SGYGILVYY 387  
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DATE 19-Mar-1997 #sequence\_revision 25-Apr-1997 #text\_change  
09-Sep-1997

ACCESSIONS S72693  
REFERENCE Smith, D.R.; Robison, K.  
#authors submitted to the EMBL Data Library, November 1993  
#submission Mycobacterium leprae cosmid B1496.  
#description #accession S72755  
#status preliminary  
##molecule\_type DNA  
##residues 1-91 ##label SMI  
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GENETICS  
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SUMMARY #length 91 #molecular-weight 9561 #checksum 2921

Query Match 74.6%; Score 47; DB 2; Length 91;  
Best Local Similarity 66.7%; Pred. No. 3.71e+01;  
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Db 56 AGIGVLSAI 64  
QY 2 AGIGILTVI 10

RESULT 13  
ENTRY A33351 #type complete  
H+-transporting ATP synthase (EC 3.6.1.34) proteolipid chain  
TITLE - *Sulfolobus acidocaldarius*  
ORGANISM #formal\_name *Sulfolobus acidocaldarius*  
DATE 20-Dec-1989 #sequence\_revision 20-Dec-1989 #text\_change  
22-Jun-1999

ACCESSIONS A33351  
REFERENCE Denda, K.; Konishi, J.; Oshima, T.; Date, T.; Yoshida, M.  
#authors Denda, K.; Konishi, J.; Oshima, T.; Date, T.; Yoshida, M.  
#JBIOL Chem. (1989) 246:7119-7121  
#title A gene encoding the proteolipid subunit of *Sulfolobus acidocaldarius* ATPase complex.  
#cross-references MUID:89214142  
#accession A33351  
#status preliminary  
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#cross-references GB:J04740; NID:9152922; PID:9152923  
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CLASSIFICATION #superfamily H+-transporting ATP synthase lipid binding  
protein hydrolase  
KEYWORDS hydrolase  
SUMMARY #length 101 #molecular-weight 10362 #checksum 4300

Query Match 74.6%; Score 47; DB 2; Length 101;  
Best Local Similarity 87.5%; Pred. No. 3.71e+01;  
Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Db 59 AAGIGVLT 66  
QY 1 AAGIGILT 8

RESULT 14  
ENTRY C70959 #type complete  
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(strain H37RV)  
ORGANISM #formal\_name *Mycobacterium tuberculosis*  
DATE 17-Jul-1998 #sequence\_revision 17-Jul-1998 #text\_change  
26-Aug-1999

ACCESSIONS C70959  
REFERENCE A70500  
#authors Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.V.; Eiglmeier, K.; Gas, S.; Barry, III, C.E.; Tekala, F.; Badcock, K.; Basham, D.; Brown, D.; Chillingworth, T.; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Hornsby, T.; Jagels, K.; Krogh, A.; McLean, J.; Moule, S.; Murphy, L.; Oliver, S.; Osborne, J.; Quail, M.A.; Rajandream, M.A.; Rogers, J.; Rutte, S.; Seeger, K.; Skelton, S.; Squares, S.; squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.

RESULT 12  
ENTRY S72755 #type complete  
B1496\_02\_163 protein - *Mycobacterium leprae*  
TITLE #formal\_name *Mycobacterium leprae*  
ORGANISM



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protein - protein database search, using Smith-Waterman algorithm  
run on: Fri May 5 22:15:32 2000: MasPar time 3.20 Seconds  
tabular output not generated.  
74.117 Million cell updates/sec

2025 RELEASE UNDER E.O. 14176

Description: (1-10) from US09267439 .pep  
Perfect Score: 63  
Sequence: 1 AAGIGIITVI 10

דנ"ה 150

According to the above:  
RAM 130  
Gap 15

2023 RELEASE UNDER E.O. 14176

post-processing: MINIMUM Match 0% Listing first 15 summaries

“geneseq” 1:geneseq

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

IMAGES

Result No.	Query Score	Match Length	DB ID	Description	Pred. No.
1	63	100.0	10	W98934 Human leukocyte antigen	4.21e+00
2	63	100.0	10	MART-1 melanoma antigen	4.21e+00
3	63	100.0	10	MART-1 epitope recogni	4.21e+00
4	63	100.0	21	W00903 Human melanoma MART-1/	4.21e+00
5	63	100.0	118	W83134 Human tumour rejection	4.21e+00
6	63	100.0	118	R63158 Tumour rejection antigen	4.21e+00
7	63	118.1	118	R84212 MART-1 melanoma antigen	4.21e+00
8	56	88.9	9	Y01751 Exemplary antigenic peptide	2.38e+01
9	56	88.9	9	Y00713 Tumour antigen booster	2.38e+01
10	56	88.9	9	Y10444 HLA Class I motif pept	2.38e+01
11	56	88.9	9	Y10567 HLA Class I motif pept	2.38e+01
12	56	88.9	9	Y10601 HLA Class I motif pept	2.38e+01
13	56	88.9	9	W54602 Peptide 1 from Melan-A	2.38e+01
14	56	88.9	9	W07379 MART-1 epitope recogni	2.38e+01
15	56	88.9	9	W77123 MART-1/MelanA syntheti	2.38e+01
16	56	88.9	9	W68380 Human MELAN-A pe	2.38e+01
17	56	88.9	9	W39430 Human immunogenic T ce	2.38e+01
18	56	88.9	9	R84196 MART-1 melanoma antigen	2.38e+01
19	56	88.9	9	W89938 Human leukocyte antigen	2.38e+01
20	56	88.9	9	W35512 MART-1/Melan A protein	2.38e+01
21	56	88.9	9	W42523 Melan A/MART epitope (	2.38e+01
22	56	88.9	10	W89839 Human leukocyte antigen	2.38e+01
23	56	88.9	10	W01550 Human leukocyte antigen	2.38e+01

The present invention describes peptides which bind to an HLA-A2 molecule and have Val at the carboxy terminus, and either: (a) Ala, Tyr or Phe at the amino terminus, and Ala at position 2 (P1); or (b) Glu at the amino terminus, and Ala, Leu, or Met at positions 2 and 3, with the proviso that Ala is not at both positions (P2). The present sequence represents an HLA-A2 binding peptide. The peptides of the present invention are used to identify HLA-A2 positive cells, provoke T cells, and determine the presence of particular T cells including cytolytic T cells (CRUS). They provide a better target than the prior art CTL-stimulating peptide.

KW metastatic melanoma; tumour-associated antigen;  
 KW immunogenic peptide; diagnosis; prognosis; prophylaxis;  
 KW therapy; vaccine.  
 OS Synthetic.  
 PN WO929193-A2.  
 PR 02-NOV-1995.  
 PR 21-APR-1995; 005063.  
 PR 22-APR-1994; US-231565.  
 PR 05-APR-1995; US-417174.  
 PR (USSH ) US SEC DEPT HEALTH .  
 PI Kawakami Y, Rosenberg SA;  
 DR WPI; 95-382963/49.  
 PT DNA encoding melanoma antigens recognised by T-lymphocytes - also  
 vectors, host cells and antibodies, used to detect, treat and  
 PT immunise animal, against melanoma.  
 PS Claim 12; Page 122; 184pp; English.  
 CC Immunogenic peptide M10-4 is a derivative of Peptide M9-2 (R84196),  
 CC which is based on the melanoma antigen (MART-1) (see R84212).  
 CC M9-2 may be modified to improve immunogenicity (see R84183-R84800)  
 CC and used in medicaments for the treatment or prevention (by  
 CC immunization) of melanoma. Antibodies against MART-1 and its  
 CC immunogenic peptides may be used in the detection and isolation of  
 CC MART-1 from a sample, the detection of which is indicative of a  
 CC disease state (melanoma or metastatic melanoma).  
 CC See also R84196.

SQ Sequence 10 AA;

Query Match 3 standard; Peptide; 10 AA.  
 ID W07381; Pred. No. 4.21e+00;  
 AC W07381; Best Local Similarity 100.0%;  
 DT 28-JUL-1997 (first entry)  
 DE MART-1 epitope recognised by melanoma specific T cell receptor.  
 KW T cell; receptor; lymphocyte; alpha; beta chain; V; variable;  
 KW J; joining; D; domain; gene segment; probe; detection;  
 KW recombination; melanoma; cancer; neoplasia; tumour; diagnosis;  
 KW MART; Melanoma Antigen Recognised by T lymphocyte.  
 OS Homo sapiens.  
 PN WO9610516-A1.  
 PD 03-OCT-1996.

PR 27-MAR-1996; U04143.  
 PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.

PJ Hwu P, Nishimura M, Rosenberg SA;

DR WPI; 96-485449/48.

PT T cell receptor alpha and/or beta chains, and related nucleic acids  
 PT - useful in pharmaceutical compsns. to prevent or treat cancer,  
 PT partic. lung, melanoma, ovarian, colon, brain or kidney tumours  
 PS Example 3; Page 11; 125pp; English  
 CC W07378-W07381 are MART-1 epitopes, M9-1, M9-2, M10-3 and M10-4  
 CC respectively, that are recognised by melanoma specific T lymphocyte  
 CC receptors (TCRs). Melanoma-specific TCRs comprising an alpha and  
 CC beta chain were made. Nucleic acids from either of these chains can be  
 CC used as probes for the detection of rearranged genes  
 CC encoding tumour associated antigens. The nucleic acids may also be used  
 CC to create transgenic animals, useful as biological models to study cancer  
 CC and evaluate diagnostic and therapeutic methods for the treatment of  
 CC cancers, particularly melanomas. Antibodies (Abs) may be raised against  
 CC alpha and beta chain polypeptides and used to detect native or denatured  
 CC TCRs and/or alterations in expression levels of T cells carrying  
 CC melanoma-specific TCRs. Abs can also purify and enrich T cells carrying  
 CC the above receptors, which can then be administered therapeutically to  
 CC mammals. Anti-idiotypic antibodies can be used to assess the level of a  
 CC specific T cell carrying these receptors in a mammal being treated using  
 CC these methods. Host cells and vectors carrying nucleic acid encoding

CC a TCR (or individual alpha or beta chain fragment) are useful in  
 CC pharmaceutical compositions to prevent or treat cancer in a mammal, e.g.  
 CC lung, melanoma, ovarian, colon, brain or kidney tumours.  
 SQ Sequence 10 AA;  
 Query Match 4 standard; Peptide; 21 AA.  
 ID WO0903;  
 AC W00903;  
 DT 23-MAY-1997 (first entry)  
 DE Human melanoma MART-1/Aa tumour associated antigen p27-47.  
 KW Adeno-associated virus; vector; liposome; transfection;  
 KW dendritic cell; melanoma; MART-1/Aa; adoptive immunotherapy;  
 KW tumour associated antigen.  
 OS Homo sapiens.  
 PN WO9703703-A1.  
 PD 06-FEB-1997; U12012.  
 PR 19-JUL-1996; US-001312.  
 PR 21-JUL-1995; US-001312.  
 PS Example 5; Page 58; 134pp; English.  
 CC Tumour associated antigens (W13660-61, W00878-903) can be loaded  
 CC into dendritic cells and used to induce antitumour immunity.  
 CC Alternatively, the dendritic cells are transfected with adeno  
 CC associated virus plasmid DNA (which includes DNA encoding the  
 CC tumour associated antigen) complexed with cationic liposomes. The  
 CC antigen loaded or transfected dendritic cells can be used to  
 CC generate tumour antigen-specific cytotoxic T lymphocytes for use in  
 CC adoptive immunotherapy in a patient having the corresponding  
 CC tumour. A suitable antigen comprises amino acids 27-47 (W00903)  
 CC of human melanoma MART-1/Aa.  
 SQ Sequence 21 AA;

RESULT 4 standard; Peptide; 21 AA.  
 ID W00878-903;  
 AC W00878-903;  
 DT 01-NOV-1995; US-007184.  
 DE MART-1 tumour associated antigen p27-47.  
 KW (RHION ) RHONE POULENC RORER PHARM INC.  
 PI Lebkowski JS, Phillip R;  
 DR WPI; 97-145208/13.  
 PT Adeno-associated virus:liposome complexes for transfecting dendritic  
 PT cells - for inducing immune response, useful for treating e.g.  
 PT neoplasia or infections  
 PS Example 5; Page 58; 134pp; English.  
 CC Tumour associated antigens (W13660-61, W00878-903) can be loaded  
 CC into dendritic cells and used to induce antitumour immunity.  
 CC Alternatively, the dendritic cells are transfected with adeno  
 CC associated virus plasmid DNA (which includes DNA encoding the  
 CC tumour associated antigen) complexed with cationic liposomes. The  
 CC antigen loaded or transfected dendritic cells can be used to  
 CC generate tumour antigen-specific cytotoxic T lymphocytes for use in  
 CC adoptive immunotherapy in a patient having the corresponding  
 CC tumour. A suitable antigen comprises amino acids 27-47 (W00903)  
 CC of human melanoma MART-1/Aa.

RESULT 5 standard; Protein; 118 AA.  
 ID W0B1314;  
 AC W0B1314;  
 DT 04-FEB-1999 (first entry)  
 DE Human tumour rejection antigen precursor.  
 KW TRAP; HIA; cancer; melanoma.  
 OS Homo sapiens.  
 PT Location/Qualifiers  
 FT Key  
 FT Misc\_difference 2 /note= "encoded by CGA"  
 FT Misc\_difference 17 /note= "encoded by GAC"  
 FT US5537476-A.  
 PN 17-NOV-1998.  
 PD 07966.

PR	03-MAR-1995; US-398409.	ID	R84212 standard; Protein; 118 AA.
PR	16-JAN-1998; US-007966.	AC	R84212;
PA	(LUW-) LUDWIG INST CANCER RES.	DT	20-APR-1996 (first entry)
PI	Boon-Falleur T, Brichard V, De Plaen E, Traversari C,	DE	MART-1 melanoma antigen.
PI	Van Pel A, Woelfelt;	KW	MART-1; melanoma antigen recognised by T-cell; melanoma;
DR	WPI; 99-043967/04.	KW	metastatic melanoma; tumour-associated antigen; immunogen;
N-PADB:	V70150.	KW	diagnosis; prognosis; prophylaxis; therapy; vaccine.
PT	Use of a tumour rejection antigen precursor - as a marker for diagnosing a disorder characterised by expression of a tumour rejection antigen precursor which is not tyrosinase	OS	Mammalian.
PT	Claim 1; Column 7-9; 11pp; English.	FH	
PS	A method has been developed for the diagnosis of a disorder which is characterised by the expression of a tumour rejection antigen precursor (TRAP) which is not tyrosinase, and which is processed to a TRA which forms a complex with an HLA-A2 molecule. The present sequence represents the TRAP for use in the present invention. The method comprises contacting a sample from a subject with an agent specific for the complex and determining the interaction between the complex and the agent as a determination of the disorder. TRAP can be used for the diagnosis and treatment of disorders characterised by the expression of the TRAP molecules such as cancers, particularly melanoma.	Key	
CC	Sequence 118 AA;	Location/Qualifiers	27..47 /note= "hydrophobic region"
CC	Query Match 6	FT	
CC	Best Local Similarity 100.0%; Score 63; DB 1; Length 118;	FT	
CC	Matches 10; Conservative 0; Pred. No. 4.21e+00; Indels 0; Gaps 0;	PN	WO9529193-A2.
Db	27 AAGIGILTVI 36	PD	02-NOV-1995
Qy	1 AAGIGILTVI 10	PF	21-APR-1994; US-221565.
SO	RESULT 6	PR	05-APR-1995; US-417174.
ID	R63158 standard; Protein; 118 AA.	PA	PA (USSH ) US SEC DEPT HEALTH.
AC	R63158;	CC	PI Kawakami Y, Rosenberg SA; WPI; 95-30963/49.
DT	26-MAY-1995 (first entry)	DR	DR N-NSDB; T02714.
DE	Tumour rejection antigen precursor.	PT	DNA encoding melanoma antigens recognised by T-lymphocytes - also vectors, host cells and antibodies, used to detect, treat and immunise animal against melanoma.
KW	Tumour rejection antigen; precursor; HLA-A2 molecule; tyrosinase;	PT	PT Claim 11; Page 117; 18pp; English.
KW	isolation; melanoma; cell line; LB-39-MEL; diagnosis; vaccine;	PS	PS Claim 11; Page 117; 18pp; English.
KW	therapy.	CC	The melanoma antigen (MART-1) is produced by recombinant DNA methods, i.e. preferably using a baculovirus vector for expression in insect cell cultures. MART-1 protein is a source of immunogenic peptides (see R84196 for peptide M9-2) which are optionally modified (see R84783-R84800) and used in medicaments for the treatment or prevention (by immunization) of melanoma. Antibodies against MART-1 and its immunogenic peptides may be used in the detection and isolation of MART-1 from a sample, the detection of which is indicative of a disease state (melanoma or metastatic melanoma).
OS	Homo sapiens.	CC	CC Sequence 118 AA;
PN	W09421126-A.	SO	SO Query Match 8
PD	29-SEP-1994.	ID	Y01751 standard; Peptide; 9 AA.
PR	18-MAR-1994; US-02487.	AC	Y01751;
PA	(LUW-) LUDWIG INST CANCER RES.	DT	25-JUN-1999 (first entry)
PI	Boon-Falleur T, Brichard V, De Plaen E, Traversari C;	DE	Exemplary antigenic peptide derived from Melan-A(MART-1).
PI	Van Pel A, Wolfel T;	KW	MAGE-3; tumour associated gene; human leucocyte antigen Class II;
DR	WPI; 94-01654/39.	KW	KW autologous CD4+ cells; MAGE-3 related disease; cancer; melanoma;
DR	Q76370.	KW	KW osteosarcoma; leukemia; carcinoma.
PT	Nucleic acid coding for a tumour rejection antigen precursor - is used for developing prods. for diagnosis or treatment of expression related disorders. partic. melanoma	OS	OS Homo sapiens.
PT	This sequence represents the tumour rejection antigen presented by HLA-A2 molecules. The tumour rejection antigen is not related to tyrosinase. The cDNA encoding this sequence was isolated from the melanoma cell line, LB-39-MEL. The tumour rejection antigen may be used for diagnosis or in vaccines or for therapy of disorders characterised by the expression of the tumour rejection antigen precursor, particularly melanoma.	PD	25-MAR-1999
PS	Sequence 118 AA;	PF	04-SEP-1998; U18601.
CC	Query Match 8	PR	04-SEP-1997; US-928615.
CC	Best Local Similarity 100.0%; Score 63; DB 1; Length 118;	PA	PA (LUDW-) LUDWIG INST CANCER RES.
CC	Matches 10; Conservative 0; Pred. No. 4.21e+00; Indels 0; Gaps 0;	CC	CC The present sequence represents an exemplary tumour associated peptide antigen. The specificity describes a MAGE-3 tumour associated gene.
Db	27 AAGIGILTVI 36	PA	PA (UVR) UNIV VRILLE BRUSSEL.
Qy	1 AAGIGILTVI 10	PI	PI Boon-Falleur T, Chaux P, Cortahals J, Heirman C, Luitjen R, Stroobant V, Thielemans K, Van Der Bruggen P;
RESULT 7	DR	DR WPI; 99-244031/20.	
PT	Isolated peptides that bind to human leucocyte antigen class II molecules	PT	PT Disclosure; Page 29; 88pp; English.
PS	Query Match 7	PS	PS The present sequence represents an exemplary tumour associated peptide antigen. The specificity describes a MAGE-3 tumour associated gene.
CC	Best Local Similarity 100.0%; Score 63; DB 1; Length 118;	CC	CC Peptides (Y01721-25) that bind human leucocyte antigen (HLA) Class I
CC	Matches 10; Conservative 0; Pred. No. 4.21e+00; Indels 0; Gaps 0;	CC	CC molecules can be derived from the MAGE-3 protein. These peptides and CC autologous CD4+ cells that bind to a complex of MAGE-3 peptide CC and HLA Class II, are used to treat MAGE-3 related diseases, CC particularly cancers (e.g. melanoma, osteosarcoma, leukemia and CC various forms of carcinoma). The peptides are also used to produce

CC specific antibodies. Detection of the peptides, e.g. in binding assays, particularly with antibodies, is used for diagnosis of such diseases.

SQ Sequence 9 AA;

Query Match Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 2.38e+01;

Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AAGIGILTV 9  
| | | | | | | |  
Qy 1 AAGIGILTV 9

RESULT 9 standard; peptide: 9 AA.  
ID Y00713; AC Y00713;  
DT 12-MAY-1999 (first entry)  
DE Tumour antigen booster peptide Melan-ANART-1 HLA-A2 #2.  
KW booster peptide; immune response modulation; allergy;  
immune response enhancer; tumour cell; tumour rejection antigen;  
leukocyte antigen-presenting molecule; autoimmune disease;  
KW allograft rejection.  
OS Homo sapiens.  
PN WO9858956-A2.  
PD 30-DEC-1998.  
PR 19-JUN-1998; U12894.  
PA (LUDWIG INSTITUTE CANCER RESEARCH)  
PI Boon-Falleur T, Uttenhoff C, Warnier G;  
DR WPI; 99-105612/09.  
PT Immunization methods using viruses expressing antigen for priming  
and booster immunizations - useful for modulating immune responses  
against antigen, e.g. enhancing immune response against tumour cells  
PT expressing tumour rejection antigens.  
PS Disclosure; Page 10; 33pp; English.

This sequence represents a tumour antigen booster peptide that can be used in the method of the invention. The method is for for modulating an immune response in a mammal against an antigen, and comprises:  
CC (A) inducing an immune response by: (i) administering a virus containing a nucleic acid molecule encoding the antigen or its precursor to generate an immune response; and (ii) administering at least one booster dose comprising a peptide including the antigen, in an adjuvant, in a combined amount effective to enhance the initial immune response; or  
CC (B) reducing an immune response as defined for (A) but using a non adjuvant with the peptide which includes the antigen, in an amount effective to reduce the initial immune response. Method (A) is used to enhance the immune response against tumour cells expressing tumour rejection antigens, and against pathogens in subjects having human leukocyte antigen-presenting molecules. Method (B) is used to reduce the immune response in allergy, autoimmune disease, and allograft rejection. Method (A) provides an immunisation method which, unlike prior art, is not limited by the host immune response against viral vectors.

SQ Sequence 9 AA;

Query Match Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 2.38e+01;

Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AAGIGILTV 9  
| | | | | | | |  
Qy 1 AAGIGILTV 9

CC assays, particularly with antibodies, is used for diagnosis of such diseases.

SQ Sequence 9 AA;

Query Match Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 2.38e+01;

Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AAGIGILTV 9  
| | | | | | | |  
Qy 1 AAGIGILTV 9

RESULT 9 standard; peptide: 9 AA.  
ID Y00713; AC Y00713;  
DT 12-MAY-1999 (first entry)  
DE HLA Class I motif Peptide SEQ ID NO:497.  
KW Cytoxic T-lymphocyte response; CTL antigen; lymphatic system;  
immunisation; tumour; infectious disease; immunotherapy; cancer;  
malignant melanoma; viral disease; hepatitis; AIDS.  
OS Synthetic.  
PN WO9858956-A2.

PT 21-JAN-1999; U14289.  
PR 10-JUL-1998; U14289.  
PR 10-DEC-1997; US-988320.  
PR 10-JUL-1997; CA-209815.  
PA (CTLI) CTL IMMUNOTHERAPIES CORP.

DR WPI; 99-120514/10.

CC Homo sapiens.  
PN WO99032183-A2.

PD 21-JAN-1999.  
PR 10-JUL-1998; U14289.

AC Y10444;

DT 12-MAY-1999 (first entry)

DE HLA Class I motif peptide SEQ ID NO:374.

KW Cytoxic T-lymphocyte response; CTL antigen; lymphatic system;

KW immunisation; tumour; infectious disease; immunotherapy; cancer;

KW malignant melanoma; viral disease; hepatitis; AIDS.

OS Synthetic.

PT Inducing a cytotoxic T lymphocyte response - by maintaining a level of antigen in the lymphatic system of a mammal so as to provide a sustained CTL response, used to treat, e.g. AIDS.  
PS Disclosure; Page 40; 199pp; English.  
CC The present invention describes a method of inducing and/or sustaining an immunological cytotoxic T lymphocyte (CTL) response in a mammal. The method comprises: (a) delivering an antigen to the mammal at a level to induce an immunological CTL response in the mammal; and (b) maintaining the level of the antigen in the mammal's lymphatic system to maintain the immunologic CTL response. The method can be used for the delivery of e.g. a differentiation antigen, a tumour-specific multilineage antigen, an embryonic antigen, an oncogene antigen, a mutated tumour-suppressor gene antigen, or a viral antigen. They can be used for the treatment of disease such as cancer, e.g. malignant melanoma or infectious disease, e.g. viral disease such as hepatitis or AIDS. Sustained antigen delivery to the lymphatic system provides for potent CTL stimulation that takes place in the milieu of the lymphoid organ, and it sustains stimulation that is necessary to keep CTL active, cytotoxic and recirculating antigens given in the present invention.

SQ Sequence 9 AA;

Query Match Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 2.38e+01;

Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AAGIGILTV 9  
| | | | | | | |  
Qy 1 AAGIGILTV 9

RESULT 11 standard; Peptide: 9 AA.  
ID Y10567; AC Y10567;

DT 12-MAY-1999 (first entry)

DE HLA Class I motif Peptide SEQ ID NO:497.

KW Cytoxic T-lymphocyte response; CTL antigen; lymphatic system;

KW immunisation; tumour; infectious disease; immunotherapy; cancer;

KW malignant melanoma; viral disease; hepatitis; AIDS.

OS Synthetic.

PN WO99032183-A2.

PD 21-JAN-1999.

PR 10-JUL-1998; U14289.

PR 10-DEC-1997; US-988320.

PR 10-JUL-1997; CA-209815.

PA (CTLI) CTL IMMUNOTHERAPIES CORP.

DR WPI; 99-120514/10.

PT Inducing a cytotoxic T lymphocyte response in a mammal. The method comprises: (a) delivering an antigen to the mammal at a level to induce an immunological CTL response in the mammal; and (b) maintaining the level of the antigen in the mammal's lymphatic system to maintain the immunologic CTL response. The method can be used for the delivery of e.g. a differentiation antigen, a tumour-specific multilineage antigen, an embryonic antigen, an oncogene antigen, a mutated tumour-suppressor gene antigen, or a viral antigen. They can be used for the treatment of disease such as cancer, e.g. malignant melanoma or infectious disease, e.g. viral disease such as hepatitis or AIDS. Sustained antigen delivery to the lymphatic system provides for potent CTL stimulation that takes place in the milieu of the lymphoid organ, and it sustains stimulation that is necessary to keep CTL active, cytotoxic and recirculating antigens given in the present invention.

CC to the lymphatic system provides for potent CTL stimulation that takes place in the milieu of the lymphoid organ, and it sustains stimulation that is necessary to keep CTL active, cytotoxic and recirculating through the body. Y10071 to Y10639 represent examples of peptide antigens given in the present invention.

SQ Sequence 9 AA;

Query Match 88.9%; Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 2.38e+01; Indels 0; Gaps 0;

Matches 9; Conservative 0; Mismatches 0; Gaps 0;

Db 1 AGIGITIV 9  
| | | | | | |  
Qy 1 AGIGITIV 9

RESULT 12  
ID Y10601 standard; Peptide; 9 AA.  
AC Y10601;  
DT 12-MAY-1999 (first entry)  
DE HUA Class I motif peptide SEQ ID NO:531.  
KW Cytotoxic T-lymphocyte response; CTL; antigen; lymphatic system;  
immunisation; tumour; infectious disease; immunotherapy; cancer;  
malignant melanoma; viral disease; hepatitis; AIDS.  
Synthetic.  
OS Homo sapiens.  
PN W09902183-A2.  
PD 21-JAN-1999.  
PF 10-JUL-1998; U14289.  
PR 10-DEC-1997; US-988320.  
RR 10-JUL-1997; CA-209815.  
PA (CTL-) CTL IMMUNOTHERAPIES CORP.  
PI Kuendig TM, Simard JU;  
DRX WPI; 99-120514/10.

PT Inducing a cytotoxic T lymphocyte response - by maintaining a level of antigen in the lymphatic system of a mammal so as to provide a sustained CTL response, used to treat, e.g. AIDS

PS Disclosure; Page 49; 199pp; English.

CC The present invention describes a method of inducing and/or sustaining an immunological cytotoxic T lymphocyte (CTL) response in a mammal. The method comprises: (a) delivering an antigen to the mammal at a level to induce an immunological CTL response in the mammal; and (b) maintaining the level of the antigen in the mammal's lymphatic system to maintain the immunologic CTL response. The method can be used for the delivery of e.g. a differentiation antigen, a tumour-specific multilneage antigen, an embryonic antigen, an oncogene antigen, a mutated tumour-suppressor gene antigen, or a viral antigen. They can be used for the treatment of disease such as cancer, e.g. malignant melanoma or infectious disease, e.g. viral disease such as hepatitis or AIDS. Sustained antigen delivery to the lymphatic system provides for potent CTL stimulation that takes place in the milieu of the lymphoid organ, and it sustains stimulation that is necessary to keep CTL active, cytotoxic and recirculating through the body. Y10071 to Y10639 represent examples of peptide antigens given in the present invention.

SQ Sequence 9 AA;

Query Match 88.9%; Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 2.38e+01; Indels 0; Gaps 0;

Matches 9; Conservative 0; Mismatches 0; Gaps 0;

Db 1 AGIGITIV 9  
| | | | | | |  
Qy 1 AGIGITIV 9

RESULT 13  
ID W54602 standard; Peptide; 9 AA.  
AC W54602;  
DT 25-SEP-1998 (first entry)  
DE Peptide 1 from Melan-A/Mart-1.  
KW Mannose; antigen-presenting cell; mannosylated peptide; T cell;  
Synthetic.

PN W09813378-A1.  
PD 02-APR-1998.  
PF 25-SEP-1997; NL0536.  
PR 26-SEP-1996; EP-202701.  
PA (UYLE-) RIDSONIV LEIDEN.  
PI Drijfhout JW, Konink F;  
WPI; 98-230631/20.  
PT Increasing uptake and presentation of antigen(s) - by adding mannose residue(s) to antigen for increasing T cell response, useful in, e.g. vaccines against viral infection(s).  
PS Disclosure; Page 24; 47pp; English.  
CC The peptides W5459-W4809 are examples of peptides to which at least 1 (preferably 2) mannose can be attached to increase their uptake as antigens by antigen-presenting cells. Uptake of agonist mannosylated peptides will increase the T cell response, whereas uptake of antagonistic peptides blocks the T cell response. Blocking binding of immunogenic autoantigens can be used in treatment of type I diabetes, rheumatoid arthritis graft rejection etc., also to induce T-cell non-responsiveness. Vaccines containing mannosylated antigen are used to prevent or treat infections by, e.g. bacteria, viruses, fungi, helminths and parasites.

SQ Sequence 9 AA;  
Query Match 88.9%; Score 56; DB 1; Length 9;  
Best Local Similarity 100.0%; Pred. No. 2.38e+01;  
Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Db 1 AAGIGITIV 9  
| | | | | | |  
Qy 1 AAGIGITIV 9

RESULT 14  
ID W07379 standard; Peptide; 9 AA.  
AC W07379;  
DT 28-JUL-1997 (first entry)  
DE MART-1 epitope recognised by melanoma specific T cell receptor.  
KW T cell; receptor; lymphocyte; alpha; beta chain; V; variable;  
KW J; joining; D; diversity; gene segment; probe; detection;  
KW recombination; melanoma; cancer; neoplasia; tumour; diagnosis;  
KW HOMO sapiens.  
OS Homo sapiens.  
PN W09630516-A1.  
PD 03-OCT-1996.

PP 27-MAR-1995; U04143.

PR 27-MAR-1995; US-411098.

PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.

PI Hwu P, Nishimura M, Rosenberg SA;

DR WPI; 96-48549/48.

PT T cell receptor alpha and/or beta chains, and related nucleic acids useful in pharmaceutical compunds. to prevent or treat cancer, particic lung, melanoma, ovarian, colon, brain or kidney tumours

PS Example 3; Page 11; 125pp; English.

CC W07379-W07381 are MART-1 epitopes, M9-1, M9-2, M10-3 and M10-4 respectively, that are recognised by melanoma specific T lymphocyte receptors (TCRs). Melanoma-specific TCRR comprising an alpha and beta chain were made. Nucleic acids from either of these chains can be used as probes for the detection of rearranged genes encoding tumour-associated antigens. The nucleic acids may also be used to create transgenic animals, useful as biological models to study cancer and evaluate diagnostic and therapeutic methods for the treatment of cancers, particularly melanomas. Antibodies (Abs) may be raised against CC TCRRs and/or alterations in expression levels of T cells carrying CC melanoma-specific TCRRs. Abs can also purify and enrich T cells carrying CC the above receptors, which can then be administered therapeutically to CC mammals. Anti-idiotypic antibodies can be used to assess the level of a CC specific T cell carrying these receptors in a mammal being treated using CC these methods. Host cells and vectors carrying nucleic acid encoding CC a TCR (or individual alpha or beta chain fragment) are useful in CC pharmaceutical compositions to prevent or treat cancer in a mammal, e.g. CC lung, melanoma, ovarian, colon, brain or kidney tumours.

SQ Sequence 9 AA;

Query Match 88.9%; Score 56; DB 1; Length 9;  
 Best Local Similarity 100.0%; Pred. No. 2.38e+01;  
 Matches 9; conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Db 1 AAGIGILTV 9  
 Qy 1 AAGIGILTV 9

## RESULT 15

ID W77123 standard; peptide: 9 AA.  
 AC W77123;  
 DT 16-NOV-1998 (first entry)  
 DE MART-1/MelanA synthetic peptide epitope 1.  
 KW Tyrosinase; tyrosinase cytotoxic lymphocyte response;  
 KW cytotoxic T lymphocyte; cysteine-depleted; melanoma.  
 OS Synthetic.  
 PN WO9833810-A2.  
 PD 06-AUG-1998.  
 PP 29-JAN-1998; U01592.  
 PR 30-JAN-1997; US-037781.

PA (UVVI-) UNIV VIRGINIA PATENT FOUND.

PI Engelhard VH, Hunt DF, Kittlesen D, Slingluff CL;  
 DR WPI; 98-137388/37.

PT Disease specific immunogen - comprises disease specific cytotoxic T lymphocyte epitope used to elicit melanoma specific CTL response  
 PS Disclosure: Page 27; 930P; English.  
 CC The peptide epitope W77119-W77138 were created for human tumour-specific cytotoxic T lymphocyte response. These peptides are are cysteine-depleted mutants of a native disease-specific CTL epitope. The cysteine-depleted CTL epitopes elicit a stronger or more specific CTL response than the native epitope. The epitopes can be used in a disease-specific immunogen to protect a mammal against disease in particular melanomas.  
 CC The peptides may also be used to screen a sample for the presence of an antigen with the same epitope, or with a different cross-reactive epitope.  
 Sequence 9 AA;

Query Match 88.9%; Score 56; DB 1; Length 9;  
 Best Local Similarity 100.0%; Pred. No. 2.38e+01;  
 Matches 9; conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Db 1 AAGIGILTV 9  
 Qy 1 AAGIGILTV 9

Search completed: Fri May 5 22:16:06 2000  
 Job time : 34 secs.

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DATE          30-Sep-1991 #sequence_revision 30-Sep-1991 #text_change
ACCESSIONS   S13592; B33358
REFERENCE    S13592
  #authors  Gardner, P.D.
  #journal Nucleic Acids Res. (1990) 18:6714
  #title   Nucleotide sequence of the epsilon-subunit of the mouse
           muscle nicotinic acetylcholine receptor.
  #cross-references MUID:91067487
  #accession S13592
  #molecule_type mRNA
  #residues  1-493 ##1label GAR
  ##cross-references EMBL:X55718; NID:g53160; PIDN:CAA39751.1; PID:953161
REFERENCE    A33358
  #authors  Buonanno, A.; Mudd, J.; Merlie, J.P.
  #journal J. Biol. Chem. (1989) 264:7611-7616
  #title   Isolation and characterization of the beta-and
           epsilon-subunit genes of mouse muscle acetylcholine
           receptor.
  #cross-references MUID:89214211
  #accession B33358
  #molecule_type DNA
  #residues  1-493 ##1label BUO
  ##cross-references GB:J04698; NID:9191599; PIDN:AAA37153.1; PID:9387086
CLASSIFICATION #superfamily acetylcholine receptor
KEYWORDS      glycoprotein; ion channel; muscle; neurotransmitter receptor;
               postsynaptic membrane; transmembrane protein
FEATURE       1-20
  #domain signal sequence #status predicted #label SIG\
  21-493   #product nicotinic acetylcholine receptor epsilon chain
            #status predicted #label MAT\
  21-239   #domain extracellular #status predicted #label EXT\
  240-266   #domain transmembrane #status predicted #label TM1\
  273-291   #domain transmembrane #status predicted #label TM2\
  307-328   #domain intracellular #status predicted #label TM3\
  329-456   #domain transmembrane #status predicted #label INT\
  457-479   #binding_site carbohydrate (Asn) (covalent) #status
  86,161-327 predicted\
  148-162   #disulfide_bonds #status predicted
SUMMARY       #length 493 #molecular_weight 54914 #checksum 1794
Query Match   Score 43; DB 1; Length 493;
  Best Local Similarity 75.0%; Pred. No. 6.64e+01;
  Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
Db          93 AGYGILRV 100
             ||:||| |
Qy          2 AGIGILTV 9

```

Search completed: Fri May 5 22:00:34 2000  
Job time : 62 secs.

#authors Gosalbes, M.J.; Perez-Gonzalez, J.A.; Gonzalez, R.; Navarro, A.  
 #journal J. Bacteriol. (1991) 173:7705-7710  
 #title Two beta-glycanase genes are clustered in *Bacillus polymyxa*:  
 molecular cloning, expression, and sequence analysis of  
 genes encoding a xylanase and an endo-beta-(1,3)-(1,  
 4)-glucanases.

#cross-references MUID:92041687  
 #accession S19011  
 #status preliminary  
 ##molecule\_type DNA  
 ##residues 1-635 ##label GOS  
 #cross-references EMBL:X57094; PID:948815; Pred. No. 4.19e+01;  
 #note the authors translated the codon GAA for residue 78 as  
 Gly, CCT for residue 272 as Thr, ATC for residue 412  
 as Glu, and ATC for residue 478 as Tyr

FUNCTION #description catalyzes the hydrolysis of 1,4-beta-xylidosic linkages in  
 xylans  
 #superfamily Clostridium xylanase A repeat homology  
 glycosidase; hydrolase; polysaccharide degradation

CLASSIFICATION FEATURE #domain Clostridium xylanase A repeat homology #label  
 408-502 CXA

SUMMARY #length 635 #molecular\_weight 67914 #checksum 2077  
 Query Match Score 44; DB 2; Length 635;  
 Best Local Similarity 75.08; Pred. No. 4.19e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Db 149 GAGIGVLT 156  
 QY :|||:|||:  
 QY 1 AAGIGILT 8

RESULT 13  
 ENTRY S01903 #type complete  
 TITLE H+-transporting ATP synthase (EC 3.6.1.34) epsilon chain -  
 ORGANISM Arabidopsis thaliana chloroplast  
 #common\_name  
 DATE 01-Dec-1989 #sequence\_revision 01-Dec-1989 #text\_change  
 22-Jun-1999  
 ACCESSIONS S01903  
 REFERENCE S01903  
 #authors Chen, H.C.; Wintz, H.; Weil, J.H.; Pillay, D.T.N.  
 #journal Nucleic Acids Res. (1988) 16:10372  
 #title Nucleotide sequence of chloroplast CF1-ATPase epsilon-subunit  
 and elongator tRNA(Met) genes from *Arabidopsis thaliana*.  
 mouse-ear cress

#cross-references MUID:89057486  
 #accession S01903  
 #molecule\_type DNA  
 #residues 1-132 ##label CHE  
 #cross-references EMBL:X12889; PID:911332; Pred. No. 6.4e+01;  
 GENETICS atpE  
 #gene chloroplast  
 #genome superfamily H+-transporting ATP synthase epsilon chain  
 CLASSIFICATION REYNDS ATP biosynthesis; chloroplast; hydrolase; membrane-associated  
 #title complex; thylakoid  
 #length 132 #molecular\_weight 14472 #checksum 1607  
 Query Match Score 43; DB 2; Length 132;  
 Best Local Similarity 66.78; Pred. No. 6.4e+01;  
 Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Db 43 AVDIGITI 51  
 QY | :|||:  
 QY 1 AAGIGILTV 9

RESULT 14  
 #type complete  
 #common\_name house mouse

ENTRY D70073 #type complete  
 TITLE metabolite transport protein homolog YXCC - *Bacillus subtilis*  
 ORGANISM #formal\_name *Bacillus subtilis*  
 DATE 05-Dec-1997 #sequence\_revision 05-Dec-1997 #text\_change  
 24-Sep-1999  
 ACCESSIONS D70073  
 REFERENCE A69580  
 #authors Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Bortero, M.G.; Bessières, P.; Brans, S.; Boulton, A.; Borchert, S.; Boriss, R.; Bourstier, L.; Brans, A.; Braun, M.; Brignell, S.C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capuano, V.; Carter, N.M.; Choi, S.K.; Codani, J.J.; Connerton, I.F.; Cummings, N.M.; Daniel, R.A.; Denizot, F.; Devine, K.M.; Duesterhoeft, A.; Ehrlich, S.D.; Emerson, P.T.; Entian, K.D.; Errington, J.; Fabre, C.; Ferrari, E.; Foulger, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Galizzi, A.; Galleron, N.; Ghim, S.Y.; Glaser, P.; Goffeau, A.; Golightly, E.J.; Grandi, G.; Guiseppi, G.; Guy, B.J.; Haiech, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Hollsappel, S.; Hosono, S.; Hull, M.F.; Itaya, M.; Jones, L.; Joris, B.; Karanata, D.; Kasahara, Y.; Klaerr-Blanchard, M.; Klein, C.; Kobayashi, Y.; Koetter, P.; Koningsstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois, S.; Lauber, J.J.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Mauel, C.; Medigue, C.; Medina, N.; Mellado, R.P.; Mizuno, M.; Moestl, D.; Nakai, S.; Noback, M.; Noone, D.; O'Reilly, M.; Ogawa, K.; Ogihara, A.; Oudega, B.; Park, S.H.; Parro, V.; Pohl, T.M.; Portetelle, D.; Porwollik, S.; Prescott, G.; A.M.; Presecan, E.; Pujic, P.; Pur nelle, B.; Rapoport, G.; Rey, M.; Reynolds, S.; Rieger, M.; Rivolta, C.; Rocha, E.; Roche, B.; Rose, M.; Sade, Y.; Sato, T.; Scanlon, E.; Schleicher, S.; Schroeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Seror, S.J.; Serron, P.; Shin, B.S.; Soldo, B.; Sorokin, A.; Tacconi, E.; Takagi, T.; Takahashi, H.; Takemaru, K.; Takeuchi, M.; Tamakoshi, A.; Tanaka, T.; Terpstra, P.; Tognoni, A.; Uchiyama, S.; Vandebroek, M.; Vanner, F.; Vassarotti, A.; Viari, A.; Veldkamp, R.; Wedler, H.; Weltzenegger, T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yanane, K.; Yasumoto, K.; Yata, K.; Yoshida, K.; Yoshikawa, H.F.; Zumstein, E.; Yoshikawa, H.; Danchin, A.

Nature (1997) 390:249-256  
 The complete genome sequence of the Gram-positive bacterium *Bacillus subtilis*.  
 #cross-references MUID:9804033  
 #accession D70073  
 #status preliminary; nucleic acid sequence not shown;  
 #molecule\_type DNA translation not shown  
 #molecule\_type DNA  
 #residues 1-461 ##label KUN  
 ##cross-references GB:299124; GB:AL009126; PID:el184706; PID:92636527  
 ##experimental\_source strain 168

GENETICS #gene YXCC  
 CLASSIFICATION #length 461 #molecular\_weight 50140 #checksum 8642  
 #summary Query Match Score 43; DB 2; Length 461;  
 Best Local Similarity 85.7%; pred. No. 6.64e+01;  
 Matches 6; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Db 285 GIGILNV 291  
 | :|||:  
 QY 3 GIGILTV 9

RESULT 15  
 #type complete  
 #common\_name house mouse  
 #formal\_name Mus musculus  
 ORGANISM

rats after partial hepatectomy and during sepsis.  
 #cross-references MUID:96003917  
 #accession S59131  
 #status preliminary  
 #molecule\_type mRNA  
 #residues 1-420 #label FUR  
 ##cross-references EMBL:D41964; NID:9604901; PID:d1008487; #checksum 4868  
**SUMMARY:** #length 420 #molecular-weight 46496 #checksum 4868  
 Query Match 80 4%; Score 45; DB 2; Length 420;  
 Best Local Similarity 55.6%; Pred. No. 2.62e+01;  
 Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;  
 Db 226 GPGYGIISV 234  
 :||:||:||:  
 Qy 1 AGIGILTV 9

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RESULT 9 D72106 #type complete  
 ENTRY #hypothetical protein - Chlamydial pneumoniae (strain CWL029)  
 TITLE #formal\_name Chlamydial pneumoniae  
 ORGANISM 23-Apr-1999 #sequence\_revision 23-Apr-1999 #text\_change  
 DATE D72106  
 A72000  
 #authors Kalmann, S.; Mitchell, W.; Marathe, R.; Lammel, C.; Fan, J.;  
 Olinger, L.; Grimwood, J.; Davis, R.W.; Stephens, R.S.  
 #journal Nature Genet. (1999) 21:385-389  
 #title Comparative genomes of Chlamydial pneumoniae and C.  
 trachomatis.  
 #cross-references MUID:992106506  
 #accession D72106  
 #status preliminary  
 #molecule\_type DNA  
 #residues 1-98 #label ARN  
 ##cross-references GB:AE001607; GB:AE001363; NID:94376474; PID:94376483  
 ##experimental\_source strain CWL029  
 GENETICS CPn0211  
 #length 98 #molecular-weight 10280 #checksum 832  
 Query Match 78 6%; Score 44; DB 2; Length 98;  
 Best Local Similarity 75.0%; Pred. No. 4.19e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 Db 54 AGVAILTV 61  
 :||:||:||:  
 Qy 2 AGIGILTV 9

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RESULT 10 C65176 #type complete  
 ENTRY qlmG protein - Escherichia coli (strain K-12)  
 TITLE #formal\_name Escherichia coli  
 ORGANISM 12-Sep-1997 #sequence\_revision 17-Sep-1997 #text\_change  
 DATE C65176  
 A64720  
 #authors Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.;  
 Burland, V.; Riley, M.; Collado-Vides, J.; Glasner, J.D.;  
 Rose, C.K.; Mayhew, G.F.; Gregor, J.; Davis, N.W.;  
 Kirkpatrick, H.A.; Goeden, M.A.; Rose, D.J.; Mau, B.; Shao,  
 Y.  
 #journal Science (1997) 277:1453-1462  
 #title The complete genome sequence of Escherichia coli K-12.  
 #cross-references MUID:97426617  
 #status nucleic acid sequence not shown; translation not shown  
 #molecule\_type DNA  
 #residues 1-456 #label BLAT  
 ##cross-references GB:AE000450; GB:U00096; NID:91790166;  
 PIDN:AC76753.1; PID:91790168; UWGP:b3730

---

##experimental\_source strain K-12, substrate MG1655  
 GENETICS qlmG  
 #gene qlmG  
 #superfamily N-acetylglucosamine-1-phosphate  
 CLASSIFICATION uridylyltransferase  
 SUMMARY #length 456 #molecular-weight 49190 #checksum 9400  
 Query Match 78.6%; Score 44; DB 2; Length 456;  
 Best Local Similarity 75.0%; Pred. No. 4.19e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 Db 124 GGIGILTV 131  
 :||:||:||:  
 Qy 2 AGIGILTV 9

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RESULT 11 H69382 #type complete  
 ENTRY ABC transporter, ATP-binding protein homolog - Archaeoglobus fulgidus  
 TITLE #formal\_name Archaeoglobus fulgidus  
 ORGANISM 05-Dec-1997 #sequence\_revision 05-Dec-1997 #text\_change  
 DATE 28-May-1999  
 ACCESSIONS H69382  
 REFERENCE A69250  
 #authors Klenk, H.P.; Clayton, R.A.; Tomb, J.F.; White, O.; Nelson, K.E.; Ketchum, K.A.; Dodson, R.J.; Gwynn, M.; Hickey, E.K.; Peterson, J.D.; Richardson, D.L.; Keravage, A.R.; Graham, D.E.; Kyriakis, N.C.; Fleischmann, G.C.; Quackerbush, J.; Lee, N.H.; Sutton, G.G.; Gill, S.; Kirkness, E.F.; Dougherty, B.A.; McKenna, K.; Adams, M.D.; Loftus, B.; Peterkin, S.; Reich, C.I.; McNeil, L.R.; Badger, J.H.; Glodek, A.; Zhou, L.; Overbeek, R.; Gocayne, J.D.; Weidman, J.F.; McDonald, L.; Utterback, T.; Cottont, M.D.; Spriggs, T.; Ariach, P.; Kaine, B.P.; Styles, S.M.; Sadow, P.W.; D'Andrea, K.P.; Bowman, C.; Fujii, C.; Garland, S.A.; Mason, T.M.; Olsen, G.J.; Fraser, C.M.; Smith, H.O.; Woese, C.R.; Venter, J.C.  
 #journal Nature (1997) 390:364-370  
 #title The complete genome sequence of the hyperthermophilic, sulfate-reducing archaeon Archaeoglobus fulgidus.

---

##cross-references MUID:98043343  
 #accession H69382  
 #status preliminary; nucleic acid sequence not shown;  
 #molecule\_type DNA 1-620 #label KLE  
 #residues 1 #cross-references GB:AE001029; GB:AE000782; NID:92689352; PID:g2649523;  
 #cross-references TIGR:AF1064  
 #label KLE  
 #domain ATP-binding cassette homology  
 #region nucleotide-binding motif A (P-loop)  
 #length 620 #molecular-weight 71200 #checksum 5979  
 CLASSIFICATION H69382  
 #superfamily ATP-binding cassette homology  
 KEYWORDS ATP; P-loop  
 FEATURE  
 #domain ATP-binding cassette homology #label ABC  
 428 612 #region nucleotide-binding motif A (P-loop)  
 SUMMARY 445 452 #length 620 #molecular-weight 71200 #checksum 5979  
 Query Match 78.6%; Score 44; DB 2; Length 620;  
 Best Local Similarity 66.7%; Pred. No. 4.19e+01;  
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;  
 Db 283 ADGLGTAV 291  
 :||:||:||:  
 Qy 1 AGIGILTV 9

---

RESULT 12 S19011 #type complete  
 ENTRY endo-1,4-beta-xylanase (EC 3.2.1.8) - Bacillus polymyxa  
 TITLE #formal\_name Bacillus polymyxa  
 ORGANISM 22-Nov-1993 #sequence\_revision 10-Nov-1995 #text\_change  
 DATE 09-Apr-1998  
 ACCESSIONS S19011  
 REFERENCE S19011

Yeast (1992) 8:397-408  
 Molecular analysis of yeast chromosome II between CMD1 and  
 LYS2: the excision repair gene RAP16 located in this region  
 belongs to a novel group of double-finger proteins.  
 MUID:92327848  
 #cross-references SGD:MUID:92327848  
 #accession SGD:S25364  
 #molecule-type DNA  
 #residues 1-47 #label MAN  
 #cross-references EMBL:X66247; NID:93548; PID:93549  
 GENETICS  
 #gene SGD:YSA1  
 #cross-references SGD:S0000315; MIPS:YBR11C  
 #map\_position 2R  
 CLASSIFICATION  
 #superfamily yffh protein; mutT domain homology  
 FEATURE  
 111-145 #domain mutT domain homology #label MUTT  
 SUMMARY #length 231 #molecular-weight 26087 #checksum 4809  
 Query Match 82.1%; Score 46; DB 1; Length 231;  
 Best Local Similarity 85.7%; Pred. No. 1.63e+01;  
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 Db 79 GIGILTI 85  
 Qy 3 GIGILTV 9  
 RESULT 7 S62369 #type complete  
 ENTRY TITLE methylcobalamin--coenzyme M methyltransferase II -  
 #molecule-type DNA  
 #cross-references GB:MI2778; NID:9334049; PIDN:AAA47464.1; PID:9334050  
 CLASSIFICATION #experimental\_source strain Becker  
 #superfamily herpesvirus glycoprotein F  
 KEYWORDS glycoprotein C.  
 FEATURE  
 1-22 #domain signal sequence #status predicted #label SIG\\  
 23-479 #product glycoprotein gIII #status predicted #label GPG\\  
 40-84 169.192.220, #binding\_site carbohydrate (Asn) (covalent) #status  
 226,285,302 predicted  
 SUMMARY #length 479 #molecular-weight 51206 #checksum 1630  
 Query Match 83.9%; Score 47; DB 1; Length 479;  
 Best Local Similarity 75.0%; Pred. No. 1.00e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 Db 456 AGIGILAI 463  
 Qy 2 AGIGILTV 9  
 RESULT 6 S62369 #type complete  
 ENTRY TITLE preliminary; nucleic acid sequence not shown  
 #status preliminary; nucleic acid sequence not shown  
 #molecule-type DNA  
 #residues 1-339 #label HAR  
 ALTERNATE\_NAMES #formal\_name Methanosaarcina Barkeri  
 ORGANISM #formal\_name Methanosaarcina Barkeri  
 DATE 19-Mar-1997 #sequence\_revision 25-Apr-1997 #text\_change  
 #cross-references EMBL:X91894; NID:91107727; PID:e204100; PID:91107728  
 SUMMARY #length 339 #molecular-weight 36761 #checksum 6431  
 Query Match 80.4%; Score 45; DB 2; Length 339;  
 Best Local Similarity 75.0%; Pred. No. 2.62e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
 Db 307 AGYGLLT 314  
 Qy 2 AGGILTV 9  
 RESULT 8 S59131 #type complete  
 ENTRY TITLE Kan-1 protein - rat  
 #status Kan-1 protein - rat  
 #molecule-type DNA  
 #residues 1-231 #label MAN  
 #cross-references EMBL:X78993; NID:9476045; PID:9476067  
 REFERENCE S45927 #formal\_name Rattus norvegicus #common\_name Norway rat  
 #authors Feldmann, H.; Manhaupt, G.; Schwarze, C.; Vetter, I.  
 #submission submitted to the Protein Sequence Database, August 1994  
 #accession S45979  
 #molecule-type DNA  
 #residues 1-231 #label FE2  
 #cross-references EMBL:Z35980; NID:9536465; PID:9536466; MIPS:YBR11C  
 REFERENCE S25364 #cross-references EMBL:Z35980  
 #authors Mannhaupt, G.; Stucke, R.; Ehmle, S.; Vetter, I.; Feldmann, H.  
 #cross-references EMBL:Z35980  
 #authors

QY 1 AAGIGILTV 9

RESULT 2 #type complete

ENTRY A33351 H+-transporting ATP synthase (EC 3.6.1.34) proteolip chain

TITLE - Sulfolobus acidocaldarius

ORGANISM #formal\_name Sulfolobus acidocaldarius

DATE 20-Dec-1989 #sequence\_revision 20-Dec-1989 #text\_change

22-Jun-1998

ACCESSIONS A33351

REFERENCE A33351 #status preliminary

#molecule\_type DNA

#residues 1-101 #label DEN

#cross-references GB:J04740; NID:9152922; PID:AAA72703\_1; PID:9152925

CLASSIFICATION #superfamily H+-transporting ATP synthase lipid-binding protein hydrolase

KEYWORDS #length 101 #molecular\_weight 10362 #checksum 4300

SUMMARY

accession A33351

#status preliminary

#molecule\_type DNA

#residues 1-101 #label DEN

#cross-references GB:J04740; NID:9152922; PID:AAA72703\_1; PID:9152925

CLASSIFICATION #superfamily H+-transporting ATP synthase lipid-binding protein hydrolase

#length 101 #molecular\_weight 10362 #checksum 4300

Query Match 83.9% Score 47; DB 2; Length 101;

Best Local Similarity 87.5% Pred. No. 1.00e+01;

Matches 7; Conservative 0; Indels 0; Gaps 0;

Db 59 AAGIGVLT 66

|:|||:|||

QY 1 AAGIGILTV 8

RESULT 3 C70599 #type complete

ENTRY hypothetical protein Rv1382 - Mycobacterium tuberculosis (strain H37RV)

TITLE

ORGANISM #formal\_name Mycobacterium tuberculosis

DATE 17-Jul-1998 #sequence\_revision 17-Jul-1998 #text\_change

26-Aug-1999

ACCESSIONS A70500

REFERENCE Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.V.; Eigle, K.; Gas, S.; Barry, J.R.; C.E.; Tekla, F.; Badcock, K.; Basham, D.; Brown, D.; Chillingworth, T.; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.; Horisby, T.; Jagels, K.; Krogh, A.; McLean, J.; Moule, S.; Murphy, L.; Oliver, S.; Osborne, J.; Quail, M.A.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.

Nature (1998) 393:537-544

#journal Deciphering the biology of Mycobacterium tuberculosis from the complete genome sequence.

#cross-references NID:91295987

#accession C70599 #status preliminary; nucleic acid sequence not shown; translation not shown

#molecule\_type DNA

#residues 1-165 #label COL

#cross-references GB:281011; GB:AL123456; PID:e275153;

GENETICS PID:91621264

CLASSIFICATION Rv1382 #superfamily Mycobacterium tuberculosis hypothetical protein

SUMMARY #length 165 #molecular\_weight 18189 #checksum 5780

Query Match 83.9% Score 47; DB 2; Length 250;

Best Local Similarity 66.7% Pred. No. 1.00e+01;

Matches 6; Conservative 1; Indels 0; Gaps 0;

Db 89 TDGIGIIV 97

Query Match 83.9% Score 47; DB 2; Length 165;

Best Local Similarity 75.0% Pred. No. 1.00e+01;

Matches 6; Conservative 0; Indels 0; Gaps 0;

Db 128 AGIGILAI 135

QY |:|||:|:|

2 AGIGILTV 9

RESULT 4 A69843 #type complete

ENTRY hypothetical protein Yjba - Bacillus subtilis

TITLE

ORGANISM #formal\_name Bacillus subtilis

DATE 05-Dec-1997 #sequence\_revision 05-Dec-1997 #text\_change

24-Sep-1998

ACCESSIONS A69843

REFERENCE A69580

#authors Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alion, G.; Azevedo, V.; Bertero, M.G.; Bassiers, P.; Bolotin, A.; Borchart, S.; Boursier, L.; Brans, A.; Braun, M.; Brignell, S.C.; Bron, S.; Brouillet, S.; Bruschi, C.V.; Caldwell, B.; Capitano, V.; Carter, N.M.; Choi, S.K.; Codani, J.J.; Connerton, I.F.; Cummings, N.J.; Daniel, R.A.; Denizot, F.; Devine, K.M.; Duesterhoeft, A.; Ehrlich, S.D.; Emmerson, P.T.; Entian, K.D.; Erbrington, J.; Fabret, C.; Ferrari, E.; Fritz, C.; Fujita, M.; Fujita, Y.; Fuma, S.; Gallizzi, A.; Galleron, N.; Ghim, S.Y.; Glaser, P.; Goffeau, A.; Golightly, B.J.; Grandi, G.; Guiseppi, G.; Guy, B.J.; Haga, K.; Haesch, J.; Harwood, C.R.; Henaut, A.; Hilbert, H.; Holsappel, S.; Hosono, S.; Hull, M.F.; Itaya, M.; Jones, L.; Joris, B.; Karamata, D.; Kasahara, Y.; Klaerr-Blanchard, M.; Klein, C.; Kobayashi, Y.; Koettter, P.; Konigstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardinois, S.; Lauber, J.; Masuda, S.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Mizuno, S.; Mauel, C.; Medigue, C.; Medina, N.; Mellado, R.P.; Mizuno, M.; Moestl, D.; Nakai, S.; Noback, M.; Noone, D.; O'Reilly, V.; Pohl, T.M.; Portebeille, D.; Porwolik, S.; Prescott, A.M.; Presecan, E.; Puig, P.; Purnelle, B.; Rapoport, G.; Rey, M.; Reynolds, S.; Rieger, M.; Rivolta, C.; Rocha, E.; Roche, B.; Rose, M.; Sadaie, Y.; Satoh, T.; Scanlon, E.; Schleicher, S.; Schoeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Seror, S.J.; Serrr, P.; Shin, B.S.; Soldo, B.; Sorokin, A.; Tacconi, E.; Takagi, T.; Takahashi, H.; Takemaru, K.; Takeuchi, M.; Tamakoshi, A.; Tanaka, T.; Terpstra, P.; Toomre, A.; Tosato, V.; Uchiyama, S.; Vandembol, M.; Vanquier, F.; Vassarotti, A.; Viari, A.; Wanbutt, R.; Wedler, B.; Weitzenecker, T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yates, K.; Yoshida, K.; Yoshikawa, H.F.; Zumstein, E.; Yoshioka, H.; Danchin, A.

Nature (1997) 390:249-256

#journal The complete genome sequence of the Gram-positive bacterium Bacillus subtilis.

#cross-references NID:98044033

#accession A69843 #status preliminary; nucleic acid sequence not shown; translation not shown

#molecule\_type DNA

#residues 1-250 #label RUN

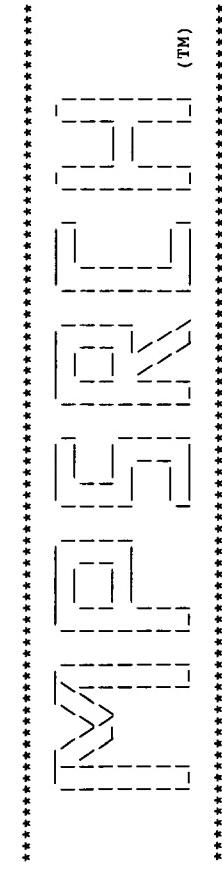
#cross-references GB:Z99110; GB:AL009126; PID:g2633472; PID:g633455

GENETICS

#gene Yjba

SUMMARY #length 250 #molecular\_weight 30119 #checksum 5271





Release 3.1a John F. Collins, Biocomputing Research Unit.

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MPsrch\_pp protein - protein database search, using Smith-Waterman algorithm

Run on: Fri May 5 21:46:46 2000; MasPar time 6.10 Seconds

577.758 Million cell updates/sec

Tabular output not generated.

Title:

Description: (1-118) from US09267439.pep

Perfect Score: 889

Sequence: 1 MRPDAHFTIYGPKKGHGS . . . . . NAPPAVEKLSAEQSQQPPYSP 118

Scoring table: PAM 150

Gap 11

Searched: 82229 seqs, 29864866 residues

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database: swiss-prot38

1:swissprot

Statistics: Mean 41.234; Variance 64.540; scale 0.639

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No. Score Query Match Length DB ID Description Pred. No.

Result No.	Score	Query	Match	Length	DB	ID	Description	Pred. No.
1	889	100.0	118	1	MAR1_HUMAN	MELANOMA ANTIGEN RECOG	2.4e-198	
2	99	11.1	344	1	CD2_MOUSE	T-CELL SURFACE ANTIGEN	4.28e-03	
3	90	10.1	628	1	LO_HUMAN	LUTHERAN BLOOD GROUP G	1.28e-01	
4	88	9.9	942	1	TMK1_ARATH	PUTATIVE RECEPTOR PROT	2.64e-01	
5	87	9.8	519	1	TYR2_HUMAN	DOPACHROMONE TAUTOMERASE	3.78e-01	
6	87	9.8	1630	1	PTP1_DROME	PROTEIN-TYROSINE PHOSP	3.78e-01	
7	86	9.7	517	1	TYR2_MOUSE	DOPACHROMONE TAUTOMERASE	5.39e-01	
8	86	9.7	704	1	MEPRB_MOUSE	A BETA-SUBUNIT	5.39e-01	
9	86	9.7	774	1	HN4G_HUMAN	HEPATOCYTE NUCLEAR FAC	5.39e-01	
10	85	9.6	320	1	PMAL_YEAST	HYPOTHETICAL	3.6e-01	
11	85	9.6	918	1	MEPRIN_A_BETA	MEMBRANE ATPASE	7.66e-01	
12	85	9.6	918	1	MEPRIN_A_BETA	SURFACE GLYCOP	1.09e+00	
13	84	9.4	327	1	CD1A_HUMAN	T-CELL SURFACE GLYCOP	1.09e+00	
14	84	9.4	334	1	Y472_RICPR	HYPOTHETICAL PROTEIN R	1.09e+00	
15	83	9.3	306	1	CDB9_MOUSE	T LYMPHOCTYE ACTIVATIO	1.53e+00	
16	83	9.3	348	1	SK1_MOUSE	SKI ONCOGENE (C-SKI)	1.53e+00	
17	83	9.3	700	1	MEPRIN_B_HUMAN	MEPRIN A BETA-SUBUNIT	1.53e+00	
18	82	9.2	426	1	UCR2_SCOPD	UBIQUITINOL-CYTOCROME C	2.16e+00	
19	82	9.2	477	1	INGR_MOUSE	INTERFERON-GAMMA RECEP	2.16e+00	
20	82	9.2	499	1	ANSP_ECOLI	L-ASPARAGINE PERMSE	2.16e+00	
21	82	9.2	897	1	CYRB_HUMAN	CYTOKINE RECEPTOR COMM	2.16e+00	
22	82	9.2	1010	1	SNI2_YEAST	SNI2 PROTEIN (SR077 PR	2.16e+00	
23	82	9.2	1013	1	PTPXP_MACNE	PROTEIN-TYROSINE PHOSP	2.16e+00	

#### ALIGNMENTS

RESULT	1	MARI_HUMAN	STANDARD;	PRT;	118 AA.
ID		O16655;			
AC		Q16655;			
DT		01-NOV-1997	(Rel. 35, Created)		
DT		01-NOV-1997	(Rel. 35, Last sequence update)		
DT		15-JUL-1998	(Rel. 36, Last annotation update)		
DE		MELANOMA ANTIGEN RECOGNIZED BY T-CELLS 1 (MART-1)	(MELAN-A PROTEIN)		
DE		(ANTIGEN SK29 AA)	(ANTIGEN LB39 AA)		
GN		MELANA OR MART1.			
OS		Homo sapiens (Human).			
OC		Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
RN	[1]				
RP		SEQUENCE FROM N.A.			
RC		TISSUE-MELANOMA;			
RX		MEDLINE; 94224770.			
RA		KAWAKAMI Y., ELIYAHU S., DELGADO C.H., ROBBINS P.F., RIVOLTINI L., TOPALIAN S.L., MIKI T., ROSENBERG S.A.;			
RA		"Cloning of the gene coding for a shared human melanoma antigen recognized by autologous T cells infiltrating into tumor."			
RT		Proc. Natl. Acad. Sci. U.S.A. 91:3515-3519(1994).			
RL		[2]			
RN		SEQUENCE FROM N.A.			
RX		MEMLINE; 94275389.			
RA		COULIE P.G., BRICHARD V., VAN PEL A., WOELFEL T., SCHNEIDER J., TRAVERSARI C., MATTEI S., DE PLAEN E., LORQUIN C., SIIKORA J.-P., RENAUD J.-C., BOON T.;			
RA		"A new gene coding for a differentiation antigen recognized by autologous cytolytic T lymphocytes on HLA-A2 melanomas."			
RT		J. Exp. Med. 180:35-42(1994).			
CC		- I - TISSUE SPECIFICITY: EXPRESSION IS RESTRICTED TO MELANOMA AND MELANOCTYE CELL LINES AND RETINA.			
CC		CC This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (see <a href="http://www.isb-sib.ch/announce/">http://www.isb-sib.ch/announce/</a> or send an email to license@isb-sib.ch).			
CC		DR EMBL; U06452; AAA19238.1; DR U06654; AAA20389.1; KW Antigen; Transmembrane. RT POTENTIAL. FT TRANSMEM 27 47. SQ SEQUENCE 118 AA; 13157 MW; DFE2CF6 CRC32;			

Query Match	100.0%	Score 889; DB 1; Length 118;
Best Local Similarity	100.0%	Pred. No. 2.48e-198;
Matches	118;	Conservative 0; Mismatches 0; Indels 0; Gaps 0;
		Cell adhesion; Signal;
		KW Immunoglobulin domain; T-cell; Glycoprotein; Antigen; Transmembrane;
DR		PIR; B28967; B28967.
		HSSP; P08921; 1A7B.
DR		MGI; MGI:88320; CD2.
KW		T-CELL SURFACE ANTIGEN CD2.
		EXTRACELLULAR (POTENTIAL).
		POTENTIAL.
		CYTOSPLASMIC (POTENTIAL).
		IG-LIKE V-TYPE DOMAIN.
		IG-LIKE C2-TYPE DOMAIN.
		PRO-RICH.
FT	SIGNAL	1 2
FT	CHAIN	23 344
FT	DOMAIN	23 202
FT	DISULFID	204 229
FT	DOMAIN	230 344
FT	DOMAIN	23 121
FT	DOMAIN	122 202
FT	DOMAIN	276 343
FT	DISULFID	133 197
FT	CARBOHYD	140 180
FT	CARBOHYD	82 82
FT	CARBOHYD	94 94
FT	CARBOHYD	135 135
FT	CARBOHYD	166 166
FT	CONFILCT	99 99
FT	CONFILCT	128 128
FT	CONFILCT	175 175
FT	CONFILCT	191 191
FT	CONFILCT	192 192
FT	SEQUENCE	344 344
FT	AA:	38414 MW: 306486 CRC32;
FT		
RESULT	2	Score 99; DB 1; Length 344;
CD2-MOUSE	STANDARD;	PRT; 344 AA.
P08920;		Pred. No. 4.38e-03;
01-NOV-1988 (Rel. 09, Created)		Gaps
01-NOV-1988 (Rel. 09, Last sequence update)		9; Mismatches 18; Indels 4;
01-FEB-1996 (Rel. 33, Last annotation update)		
T-CELL SURFACE ANTIGEN CD2 PRECURSOR (T-CELL SURFACE ANTIGEN		
T11/LEU-5) (LFA-3 RECEPTOR).		
CD2.		
Mus musculus (Mouse).		
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;		
Eutheria; Rodentia; Sciurognathii; Muridae; Murinae; Mus.		
[1]		
SEQUENCE FROM N.A.		
STRAIN=B10.A;		
MEDLINE: 87276135.		
SEWELL W.A., BROWN M.H., FINK P.J., KOZAK C.A., CRUMPTON M.J.;		
"The murine homologue of the T lymphocyte CD2 antigen: molecular cloning, chromosome assignment and cell surface expression."		
Eur. J. Immunol. 17:1015-1020(1987). [2]		
SEQUENCE FROM N.A.		
MEDLINE: 88004738.		
CLAYTON L.K., SAYRE P.H., NOVOTNY J., REINHERZ E.L.;		
"Murine and human T11 (CD2) cDNA sequences suggest a common signal transduction mechanism".		
Eur. J. Immunol. 17:1367-1370(1987). [3]		
SEQUENCE FROM N.A.		
STRAIN=BALB/C; TISSUE=LIVER;		
MEDLINE: 88144486.		
DIAMOND D.J., CLAYTON L.K., SAYRE P.H., REINHERZ E.L.;		
"Exon-Intron organization and sequence comparison of human and murine T11 (CD2) genes".		
Proc. Natl. Acad. Sci. U.S.A. 85:1615-1619(1988).		
-1- FUNCTION: CD2 INTERACTS WITH LYMPHOCYTE FUNCTION-ASSOCIATED ANTIGEN (LFA-3) AND OX-45/BGM-1 TO MEDIATE ADHESION BETWEEN T CELLS AND OTHER CELL TYPES. CD2 IS IMPLICATED IN THE TRIGGERING OF T-CELLS. THE CYTOPLASMIC DOMAIN IS IMPLICATED IN THE SIGNALING FUNCTION.		
-1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.		
-1- SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN SUPERFAMILY.		
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EMBL; Y00023; CAA88258; -;		
EMBL; Y06143; CAA89500; -;		
EMBL; M19807; AAA37393; -;		
EMBL; M19799; AAA37393; -;		
EMBL; M19801; AAA37393; -;		
EMBL; M19803; AAA37393; -;		
EMBL; M19805; AAA37393; -;		
PIR; S02293; S02293;		
PIR; S03347; S03347;		
RL	Cancer Res. 54: 5761-5765 (1994)	
RL	Medline; 95042397.	
CC	-1- FUNCTION: PROBABLE RECEPTOR. MAY MEDIATE INTRACELLULAR SIGNALING.	
CC	-1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.	
CC	-1- TISSUE SPECIFICITY: WIDE TISSUE DISTRIBUTION (HIGHEST IN THE PANCREAS AND VERY LOW IN BRAIN). CLOSELY ASSOCIATED WITH THE BASAL LAYER OF CELLS IN EPITHELIUM AND THE ENDOTHELIUM OF BLOOD VESSEL WALLS.	
CC	"Developmental Stage: IS UNDER DEVELOPMENTAL CONTROL IN LIVER AND	

MAY ALSO BE REGULATED DURING DIFFERENTIATION IN OTHER TISSUES.  
UPREGULATED FOLLOWING MALIGNANT TRANSFORMATION IN SOME CELL TYPES.  
- 1 - POLYMORPHISM: LU IS RESPONSIBLE FOR THE LUTHERAN BLOOD GROUP  
SYSTEM.  
- 1 - SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN SUPERFAMILY. CONTAINS  
3 C2-LIKE AND 2 V-LIKE DOMAINS.

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EMBL;		X84225;	CAA56449	1;	-;
EMBL;		X80006;	CAA56327	1;	-;
MIM;		111200;	-;		
PFAM;		PFO0047;	19;	4;	
Receptor; Immunoglobulin domain; Glycoprotein; Transmembrane; Signal;					
Blood group antigen.		SIGNAL	1	31	
T	T	CHAIN	32	628	LUTHERAN BLOOD GROUP GLYCOPROTEIN.
T	T	DOMAIN	32	547	EXTRACELLULAR (POTENTIAL).
T	T	TRANSMEM	548	568	POTENTIAL.
T	T	DOMAIN	569	628	CYTOSPLASMIC (POTENTIAL).
T	T	DOMAIN	46	132	IG-LIKE V-TYPE DOMAIN.
T	T	DOMAIN	165	244	IG-LIKE V-TYPE DOMAIN.
T	T	DOMAIN	284	344	IG-LIKE C2-TYPE DOMAIN.
T	T	DOMAIN	377	431	IG-LIKE C2-TYPE DOMAIN.
T	T	DOMAIN	466	529	IG-LIKE C2-TYPE DOMAIN.
T	T	DISULFID	53	125	PROBABLE.
T	T	DISULFID	172	237	PROBABLE.
T	T	DISULFID	291	337	PROBABLE.
T	T	DISULFID	384	424	PROBABLE.
T	T	DISULFID	473	522	PROBABLE.
T	T	CARBOHYD	321	321	POTENTIAL.
T	T	CARBOHYD	377	377	POTENTIAL.
T	T	CARBOHYD	383	383	POTENTIAL.
T	T	CARBOHYD	419	419	POTENTIAL.
T	T	CARBOHYD	439	439	POTENTIAL.
T	T	CONFLICT	225	226	RL -> PC ( IN REF. 2).
T	T	CONFLICT	355	356	EL -> DV ( IN REF. 2).

2	SEQUENCE	628 AA;	67374 MW;	2D577E45 CRC32;
	Query Match	10.1%;	Score 90;	DB 1; Length 628;
	Best Local Similarity	34.4%;	Pred. No.	1.28e-01;
	Matches	11;	Mismatches	8;
	Conservative	12;	Indels	1;
			Gaps	1
3	544 TSONGVA/NAVASVGLLILWA/FYCVERKG	575		
4	: : :   :   :    :   :   :   :   :   :			
5	23 TAAEAGIGILTVLGVLILLI-GCWYCRRRNG	53		

RESULT	4	TMK1_ARATH	STANDARD;	PRT;	942 AA.
C	C	PS3298;			
F	F	01 NOV-1995	(Rel. 32, Created)		
F	F	01 NOV-1995	(Rel. 32, Last sequence update)		
F	F	15 JUL-1998	(Rel. 36, Last annotation update)		

**3** ?**PUTATIVE RECEPTOR PROTEIN KINASE TMK1 PRECURSOR (EC 2.7.1.-).**  
TMK1.  
*Arabidopsis thaliana* (Mouse-ear cress).  
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophytes;  
euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;  
core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae.  
*Arabidopsis*.

SEQUENCE FROM N.A.  
STRAINE-CV; 93076110  
CHANG C., SCHALLER G.E., PATTERSON S.E., KWOK S.F.,  
L.J.

RX MEDLINE; 94198295.  
 RA YOKOYAMA K., SUZUKI H., YASUMOTO K.I., TOMITA Y., SHIBAHARA S.;  
 RT "Molecular cloning and functional analysis of a cDNA coding for human  
 DOPACHROME tautomerase/tyrosinase-related protein-2.";  
 RP Blochin. Biophys. Acta 1217:317-321(1994).  
 RL [2]  
 RN SEQUENCE FROM N.A.  
 RP SEQUENCE N.A.  
 RX MEDLINE; 94266170.  
 RA CASSADY J.L., STURM R.A.;  
 RT "Sequence of the human dopachrome tautomerase-encoding TRP-2 cDNA.";  
 RL Gene 143:295-298(1994).  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE; 94139684.  
 RA BOICHARD B., DEL MARMOL V., JACKSON I.J., CHERIF D., DUBERTRET L.;  
 RT "Molecular characterization of a human tyrosinase-related-protein-2  
 cDNA. Patterns of expression in melanocytic cells.";  
 RL Eur. J. Biochem. 219:127-134(1994).  
 RN SEQUENCE OF 1-98 FROM N.A.  
 RC TISSUE-LIVER;  
 RX MEDLINE; 9607908.  
 RA YOKOYAMA K., O'SULLIVAN B.J., BOX N.F., SMITH A.G., SMITH S.E.,  
 RT "Cloning of the human Dopachrome tautomerase/tyrosinase-related  
 protein 2 gene and identification of two regulatory regions required  
 for its pigment cell-specific expression.";  
 RL J. Biol. Chem. 269:27080-27087(1994).  
 CC -I CATALYTIC ACTIVITY: L-DOPACHROME = 5,6-DIHYDROXYINDOLE-2-  
 CARBOXYLATE.  
 CC -I COFACTOR: BINDS TWO ZINC IONS (BY SIMILARITY).  
 CC -I PATHWAY: MELANIN BIOSYNTHESIS.  
 CC -I SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN. MELANOSOMAL.  
 CC -I SIMILARITY: BELONGS TO THE TYROSINASE FAMILY.  
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration  
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 CC -----  
 DR EMBL; D17547; BAA04484.1; PROSITE; PS00497; TYROSINASE\_1; PROSITE;  
 DR EMBL; L18967; AAA20870.1; PFAM; PF00264; tyrosinase\_1.  
 KW Isomerase; Zinc; Glycoprotein; Signal; Transmembrane;  
 KW Melanin biosynthesis.  
 FT SIGNAL 1 PIR; S43510; S43510.  
 DR MIM; 191275; PROSITE; PS00497; TYROSINASE\_1; PROSITE;  
 DR DOMAIN 24 P50498; TYROSINASE\_2; PROSITE;  
 DR DOMAIN 24 L18967; AAA20870.1; PFAM; PF00264; tyrosinase\_1.  
 FT DOMAIN 473 Isomerase; Zinc; Glycoprotein; Signal; Transmembrane;  
 FT DOMAIN 494 LUMENAL, MELANOSOME (POTENTIAL).  
 FT METAL 189 189 POTENTIAL.  
 FT CHAIN 24 519 DOPACHROME TAUTOMERASE.  
 FT DOMAIN 24 472 LUMENAL, MELANOSOME (POTENTIAL).  
 FT DOMAIN 494 519 CYTOPLASMIC (POTENTIAL).  
 FT METAL 211 211 ZINC A (BY SIMILARITY).  
 FT METAL 220 220 ZINC A (BY SIMILARITY).  
 FT METAL 369 369 ZINC B (BY SIMILARITY).  
 FT METAL 373 373 ZINC B (BY SIMILARITY).  
 SQ SEQUENCE 519 AA; 59145 MW; 4FEEFCDD2 CRC32;

Query Match 9.8%; Score 87; DB 1; Length 519;  
 Best Local Similarity 35.3%; Pred. No. 3 78e-01;  
 Matches 12; Conservative 11; Mismatches 10; Indels 1; Gaps 1;

Db 477 MGTLVALVAGLFLVLLAFLQYRRLKGTYPLMETHL 510  
 Qy 30 IGILTIVLIGVLLIGCWCRR-RNGYALMDRSL 62

**RESULT 6**  
 ID PTPL\_DROME STANDARD PRT; 1630 AA.  
 AC P25992;  
 DT 01-JUN-1994 (Rel. 29, Created)  
 DT 01-JUN-1994 (Rel. 29, Last sequence update)  
 DT 15-DEC-1999 (Rel. 39, Last annotation update)  
 DE PROTEIN-TYROSINE PHOSPHATASE 10 PRECURSOR (EC 3.1.3.48) (RECEPTOR-LINKED PROTEIN-TYROSINE PHOSPHATASE 10D).  
 DE PPI10D.  
 OS Drosophila melanogaster (Fruit fly).  
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;  
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
 OC Rhadroidea; Drosophilidae; Drosophila.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=EMBRYO;  
 RX MEDLINE; 92034989.  
 RA TIAN S.-S., TSOUFLAS P., ZINN K.;  
 RT "Three receptor-linked protein-tyrosine phosphatases are selectively expressed on central nervous system axons in the Drosophila embryo.";  
 RT Cell 67:675-685(1991).  
 RL [2]  
 RN PPI10D.  
 RP SEQUENCE FROM N.A.  
 RC TISSUE=EMBRYO;  
 RX MEDLINE; 92034988.  
 RA YANG X., SEO K.T., BAHRI S.M., OON S.H., CHIA W.;  
 RT "Two Drosophila receptor-like tyrosine phosphatase genes are expressed in a subset of developing axons and pioneer neurons in the embryonic CNS.";  
 RT Cell 67:661-673(1991).  
 CC -I CATALYTIC ACTIVITY: PROTEIN-TYROSINE PHOSPHATE + H(2)O = PROTEIN-TYROSINE + ORTHOPHOSPHATE.  
 CC -I SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.  
 CC -I ALTERNATIVE PRODUCTS: TWO ISOFORMS THAT DIFFER IN THEIR C-TERMINAL TAILS ARE PRODUCED BY ALTERNATIVE SPlicing.  
 CC -I TISSUE SPECIFICITY: SELECTIVELY EXPRESSED IN A SUBSET OF AXONS AND PIONEER NEURONS IN THE EMBRYO.  
 CC -I SIMILARITY: 1 PROTEIN-TYROSINE PHOSPHATASE DOMAIN.  
 CC -I SIMILARITY: CONTAINS 12 FIBRONECTIN TYPE III-LIKE DOMAINS.  
 CC -----  
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 CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).  
 CC -----  
 DR EMBL; M80538; AA28952.1; PROSITE; PS00497; TYROSINASE\_1; PROSITE;  
 DR EMBL; M80465; AA28484.1; PROSITE; PS00498; TYROSINASE\_2; PROSITE;  
 DR PIR; C41214; C41214; PROTEIN-TYROSINE PHOSPHATASE 10 PRECURSOR (EC 3.1.3.48);  
 DR PIR; D41214; D41214; PROTEIN-TYROSINE PHOSPHATASE 10 PRECURSOR (EC 3.1.3.48);  
 DR HSSP; P18052; LYFO; FLYBASE; FBgn0004370; Ptp10D.

PROSITE; PS00383; TYR_PHOSPHATASE_1;	
PROSITE; PS50056; TYR_PHOSPHATASE_2;	
PROSITE; PS50055; TYR_PHOSPHATASE_PTP;	
PFAM; PF00041; fn3; 10.	
PFAM; PF00102; Y_PhoPhatase;	
Transmembrane; Hydrolase; Duplication; Signal; Alternative splicing;	
Repeat.	
SIGNAL	1 34
CHAIN	35 1630
DOMAIN	35 1196
TRANSEM	1197 1630
DOMAIN	1219 1218
DOMAIN	43 119
DOMAIN	120 214
DOMAIN	215 308
DOMAIN	309 402
DOMAIN	403 493
DOMAIN	494 580
DOMAIN	581 669
DOMAIN	670 766
DOMAIN	767 861
DOMAIN	862 955
DOMAIN	956 1048
DOMAIN	1049 1189
DOMAIN	1290 1532
CARBOHYD	75 75
CARBOHYD	106 106
CARBOHYD	128 128
CARBOHYD	169 169
CARBOHYD	212 212
CARBOHYD	229 229
CARBOHYD	259 259
CARBOHYD	289 289
CARBOHYD	317 317
CARBOHYD	471 471
CARBOHYD	486 486
CARBOHYD	512 512
CARBOHYD	533 533
CARBOHYD	588 588
CARBOHYD	668 668
CARBOHYD	687 687
CARBOHYD	719 719
CARBOHYD	723 723
CARBOHYD	823 823
CARBOHYD	841 841
CARBOHYD	874 874
CARBOHYD	908 908
CARBOHYD	925 925
CARBOHYD	1001 1001
CARBOHYD	1104 1104
CARBOHYD	1135 1135
CARBOHYD	1194 1194
ACT_SITE	1467 1467
VARSPLIC	1548 1557
TRANSEM	1558 1630
TRANSEM	124 124
TRANSEM	127 127
TRANSEM	1125 1125
TRANSEM	1167 1167
TRANSEM	1172 1172
TRANSEM	1216 1216
TRANSEM	1457 1457
SEQUENCE	1630 AA; 184861 MW;
VARSPLIC	MISSING (IN SHORT ISOFORM).
TRANSEM	D -> I (IN REF. 2).
TRANSEM	S -> L (IN REF. 2).
TRANSEM	Y -> YQ (IN REF. 1).
TRANSEM	IG -> YR (IN REF. 1).
TRANSEM	R -> A (IN REF. 1).
TRANSEM	I -> L (IN REF. 1).
TRANSEM	C -> G (IN REF. 1).
SEQUENCE	GQQVQDENG -> DDEGIAESGM (IN SHORT ISOFORM).
VARSPLIC	MISSING (IN SHORT ISOFORM).
TRANSEM	DB 1; Length 1630;
TRANSEM	Pred. No. 3.78e-01;
TRANSEM	Mismatches 7; Indels 3; Gaps
*Query Match Similarity 9.8%; Score 87;	
Best Local Matches 37.0%; Pred. No. 3.78e-01;	
Matches 17; Conservative 7; Mismatches 19;	
Indels 3; Gaps	
1184 YSPPIOTD-QDNTSLIVATVPP-LTLL-VLLVLIFYKRRRNCR 1226	
10 YGPKKGHGSYTAEEAGIGILTVLGVLLIGCWYCRRLNGIR 55	

TO A DARK GREY/BROWN EUMELANIN. THE SLATEY-20 MUTATION HAS A SIMILAR PHENOTYPE, THE SLATEY-20 (LIGHT) MUTATION HAS A MORE SEVERE EFFECT AND IS SEMIDOMINANT. ITS PHENOTYPE MAY BE A RESULT OF THE FAILURE OF THE ENZYME TO BE CORRECTLY TARGETED TO ITS NORMAL LOCATION ON THE INNER FACE OF THE MELANOSOME MEMBRANE.  
-!- SIMILARITY: BELONGS TO THE TYROSINASE FAMILY.

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DR . EMBL; X63349; CAA44951.1; .  
 DR . PIR; S19243; S19243 .  
 DR . MGI; MGI:102563; DCT.  
 DR PROST1; PS00497; TYROSTINASE\_1; 1.  
 DR PROST1; PS00498; TYROSTINASE\_2; 1.  
 DR PFAM; PF00264; Tyrosinase; 1.  
 KW Isomerase; Zinc; Glycoprotein; Signal; Transmembrane;  
 KW Melanin biosynthesis; Disease mutation;  
 FT SIGNAL 1 23 POTENTIAL.  
 FT CHAIN 24 517 DOPACHROME TAUTOMERASE.  
 FT DOMAIN 24 472 LUMENAL, MELANOSOME (POTENTIAL).  
 FT TRANSMEM 473 491 POTENTIAL.

FT DOMAIN 492 517 CYTOPLASMIC (POTENTIAL).  
 FT METAL 189 189 ZINC A (BY SIMILARITY).  
 FT METAL 211 211 ZINC A (BY SIMILARITY).  
 FT METAL 220 220 ZINC A (BY SIMILARITY).  
 FT METAL 369 369 ZINC B (BY SIMILARITY).  
 FT METAL 373 373 ZINC B (BY SIMILARITY).  
 FT METAL 396 396 ZINC B (BY SIMILARITY).  
 FT CARBOHYD 92 92 POTENTIAL.  
 FT CARBOHYD 170 170 POTENTIAL.  
 FT CARBOHYD 178 178 POTENTIAL.  
 FT CARBOHYD 237 237 POTENTIAL.  
 FT CARBOHYD 300 300 POTENTIAL.  
 FT CARBOHYD 342 342 POTENTIAL.  
 FT CARBOHYD 377 377 POTENTIAL.  
 FT VARIANT 194 194 R -> Q (IN SLATY).  
 FT VARIANT 434 434 P -> L (IN SLATY-2J).  
 FT VARIANT 486 486 G -> (IN SLATY-LT).  
 SQ SEQUENCE 517 AA; 58569 MW; 8EBA041 CRC32;

Query Match 9.7% Score 86; DB 1; Length 517;

Best Local Similarity 41.28% Pred. No. 5.39e-01; Matches 14; Conservative 9; Mismatches 10; Indels 1; Gaps 1;

Db 475 IGLGAFVLLGLIAFLQYRRLRKSYAPLMETGL 508  
 | ||| : : ||| : ||| : ||| : ||| : ||| : ||| : |||  
 Qy 30 IGILTIVLGVLIGCWCYCR-RNGYRALMDKSL 62

RESULT 8  
 ID MEPB\_MOUSE STANDARD; PRT; 704 AA.  
 AC Q61847.  
 DT 01-NOV-1997 (Rel. 35, Created)  
 DT 01-NOV-1997 (Rel. 35, Last sequence update)  
 DE MEPRIN A BETA-SUBUNIT PRECURSOR (EC 3.4.24.18) (ENDOPEPTIDASE-2).  
 GN MEPB OR MEP-1B.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
 OC Eutheria; Rodentia; Sciurognathii; Muridae; Murinae; Mus  
 [1] RN SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.  
 RC TISSUE-KIDNEY;  
 RX MEDLINE; 94012651.  
 RA GORBEA C.M., MARCHAND P., JIANG W., COPELAND N.G., GILBERT D.J.,  
 RA JENKINS N.A., BOND J.S.,  
 RA "Cloning, expression, and chromosomal localization of the mouse  
 RT meprin beta subunit,";  
 RL J. Biol. Chem. 268:21035-21043 (1993).  
 RN [2] SEQUENCE FROM N.A. (ISOFORM BETA').  
 RC TISSUE-KIDNEY;  
 RX MEDLINE; 96147211.  
 RA DIETRICH J.M., BOND J.S., JIANG W.;  
 RA "A novel meprin beta' mRNA in mouse embryonal and human colon  
 carcinoma cells.";  
 RT

RL J. Biol. Chem. 271:2271-2278(1996).  
 CC 1- CATALYTIC ACTIVITY: HYDROLYSIS OF PROTEIN AND PEPTIDE SUBSTRATES  
 CC PREFERENTIALLY ON CARBOXYL SIDE OF HYDROPHOBIC RESIDUES.  
 CC -1- COFACTOR: BINDS ONE ZINC ION.  
 CC -1- SUBUNIT: HETEROOTERAMER OF TWO ALPHA AND TWO BETA SUBUNITS WHICH  
 CC IS FORMED BY THE NON-COVALENT ASSOCIATION OF TWO DISULFIDE-LINKED  
 CC HETERODIMERS.  
 CC -1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.  
 CC -1- ALTERNATIVE PRODUCTS: 2 ISOFORMS, THE BETA AND BETA' SUBUNITS.  
 CC DIFFER IN THEIR N-TERMINUS DUE TO DIFFERENTIAL PROMOTER USAGE AND  
 CC ALTERNATIVE SPlicing.  
 CC -1- TISSUE SPECIFICITY: THE BETA-SUBUNIT IS EXPRESSED IN KIDNEY,  
 CC INTESTINAL BRUSH BORDERs, AND SALIVARY DUCTS. THE BETA' -ISOFORM  
 CC HAS BEEN FOUND IN CARCINOMA CELLS.  
 CC -1- INDUCTION: THE BETA' -SUBUNIT IS INDUCED BY THE MORPHOGEN RETINOIC  
 CC ACID.  
 CC -1- PIM: THIS PROTEIN UNDERGOES PROTEOLYTIC PROCESSING. BOTH FORMS  
 CC ARE GLYCOSYLATED.  
 CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY M12A (ZINC  
 CC METALLOROTEASE); ALSO KNOWN AS THE ASTACIN SUBFAMILY.  
 CC -1- SIMILARITY: CONTAINS 1 MAM DOMAIN.  
 CC -1- SIMILARITY: CONTAINS 1 EGF-LIKE DOMAIN.  
 CC -1- This SWISS-PROT entry is copyright. It is produced through a collaboration  
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 CC or send an email to [license@lsb-sib.ch](mailto:license@lsb-sib.ch)).

CC -1- EMBL; L15193; AAA75234.1; -.  
 DR HSSP; P28825; 11AF.  
 DR MGD; MGI:96364; MEP1B.  
 DR PROSITE; PS00142; ZINC-PROTEASE; 1.  
 DR PROSITE; PS00740; MAM; 1.  
 DR PROSITE; PS00060; MAM; 2; 1.  
 DR PROSITE; PS00022; EGF; 1; FALSE\_NEG.  
 DR PROSITE; PS00186; EGF; 2; FALSE\_NEG.  
 DR PROTEAM; PF00098; EGF; 1.  
 DR PROTEAM; PF00629; MAM; 1.  
 DR PROTEAM; PF00917; MATH; 1.  
 DR PROTEAM; PF01400; Astacin; 1.  
 KW Transmembrane; Hydrolase; Metalloprotease; Zinc; Glycoprotein;  
 KW Zyrogen; Signal; EGF-like domain; Alternative splicing.  
 FT SIGNAL 1 20 POTENTIAL.  
 FT PROPEP 21 64 BY SIMILARITY.  
 FT CHAIN 65 704 MERIN A BETA-SUBUNIT.  
 FT DOMAIN 21 654 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 655 678 CYTOPLASMIC (POTENTIAL).  
 FT DOMAIN 679 704 METALLOPROTEASE.  
 FT DOMAIN 63 260 MAM.  
 FT DOMAIN 607 647 EGF-LIKE.  
 FT METAL 153 155 ZINC (CATALYTIC) (BY SIMILARITY).  
 FT ACT\_SITE 154 154 BY SIMILARITY.  
 FT METAL 157 157 ZINC (CATALYTIC) (BY SIMILARITY).  
 FT METAL 163 163 ZINC (CATALYTIC) (BY SIMILARITY).  
 FT DISULFID 611 622 BY SIMILARITY.  
 FT DISULFID 616 631 BY SIMILARITY.  
 FT DISULFID 633 646 BY SIMILARITY.  
 FT CARBOHYD 193 193 POTENTIAL.  
 FT CARBOHYD 219 219 POTENTIAL.  
 FT CARBOHYD 255 255 POTENTIAL.  
 FT CARBOHYD 316 316 POTENTIAL.  
 FT CARBOHYD 422 422 POTENTIAL.  
 FT CARBOHYD 437 437 POTENTIAL.  
 FT CARBOHYD 529 529 POTENTIAL.  
 FT CARBOHYD 548 548 POTENTIAL.  
 FT CARBOHYD 593 593 POTENTIAL.  
 FT VARSPLIT 1 27 BY SIMILARITY.  
 FT

SQ	SEQUENCE	704 AA;	79548 MW;	83CF75C1 CRC32;	Saccharomyces cerevisiae; Saccharomyces.
	Query Match	9.7%	Score 86;	DB 1; Length 704;	[1] SOURCE FROM N.A.
	Best Local Similarity	26.28;	Pred. No. 5.39e-01;		SPRAIN-S288C;
	Matches	11;	Conservative	Indels 14;	RC
			Mismatches	3;	RX
				Gaps 3;	RA
Db	648 KRGSTRDTVIIAVSSSTVYFAVML-ITLVSV-YCTRRK-YR 686				DOIGNON F., BITEAU N., CROUZET M., AIGLE M.;
Qy	14 RKGHIGHSYTAAEAGIGLTVLIGLJGWCRRNSYR 55				"The complete sequence of a 19,482 bp segment located on the right arm of chromosome II from <i>Saccharomyces cerevisiae</i> ."
					RL Yearst 9:189-199(1993).
					-1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
CC					CC
					This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <a href="http://www.isb-sib.ch/announce/">http://www.isb-sib.ch/announce/</a> or send an email to license@isb-sib.ch).
CC					CC
CC					This SWISS-PROT entry is copyright. It is produced through a collaboration between the Swiss Institute of Bioinformatics and the EMBL outstation - the European Bioinformatics Institute. There are no restrictions on its use by non-profit institutions as long as its content is in no way modified and this statement is not removed. Usage by and for commercial entities requires a license agreement (See <a href="http://www.isb-sib.ch/announce/">http://www.isb-sib.ch/announce/</a> or send an email to license@isb-sib.ch).
CC					CC
CC					DR ENBL; X70529; CAA9930.1; -.
CC					DR EMBL; 23613A; CAR85228.1; -.
CC					DR PIR; S2966; S32966.
CC					DR PFAM; PF00106; adh_short; 1.
CC					DR PIRAM; PE00106; adh_short; 1.
CC					KW Hypothetical protein; Transmembrane POTENTIAL.
CC					FT TRANSMEM 162 182 POTENTIAL.
CC					FT TRANSMEM 255 275 POTENTIAL.
CC					FT TRANSMEM 280 300 POTENTIAL.
CC					FT SEQUENCE 320 AA; 35986 MW; CRC32;
CC					Query Match 9.6%; Score 85; DB 1; Length 320;
CC					Best Local Similarity 41.7%; Pred. No. 7.66e-01;
CC					Matches 10; Conservative 8; Mismatches 4; Indels 2; Gaps 2;
CC					DR 287 FGVLNLIVPPYMGGSWYIRKW 310
CC					QY 30 IGLITVILGV-LLLIGC-WYCRRR 51
CC					RESULT 11 ID MPBP_RAT STANDARD; PPF; 704 AA.
CC					AC P28826;
CC					DT 01-DEC-1992 (Rel. 24, Created)
CC					DT 01-FEB-1996 (Rel. 33, Last sequence update)
CC					DT 15-DEC-1998 (Rel. 37, Last annotation update)
CC					DE MPBP_A BETA-SUBUNIT PRECURSOR (EC 3.4.24.18) (ENDOPEPTIDASE 2).
CC					GN MPBP.
CC					OS Rattus norvegicus (Rat).
CC					OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
CC					OC Batheria; Rodentia; Sciurognathia; Muridae; Murinae; Rattus.
CC					RN [1]
CC					RP SEQUENCE FROM N.A.; AND PARTIAL SEQUENCE.
CC					RC STRAIN=SPRAGUE-DAWLEY; TISSUE=KIDNEY;
CC					RX MEDLINE; 92317075
CC					RA JOHNSON G.D., HERSH L.B.;
CC					RT Cloning a rat mpbp cDNA reveals the enzyme is a heterodimer.
CC					RL J. Biol. Chem. 267:13305-13512(1992).
CC					RN [2]
CC					RP ERERATUM (REFRACTION).
CC					RX MEDLINE; 93359474.
CC					RA JOHNSON G.D., HERSH L.B.;
CC					RL J. Biol. Chem. 268:17447-17647(1993).
CC					-1- CATALYTIC ACTIVITY: HYDROLYSIS OF PROTEIN AND PEPTIDE SUBSTRATES PREFERENTIALLY ON CARBOXYL SIDE OF HYDROPHOBIC RESIDUES.
CC					CC -1- COFACTOR: Binds one ZINC ION.
CC					CC -1- SUBUNIT: HETERODIMER OF TWO ALPHA AND TWO BETA SUBUNITS WHICH IS FORMED BY THE NON-COVALENT ASSOCIATION OF TWO DISULFIDE-LINKED HETERODIMERS.
CC					-1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.
CC					CC -1- TISSUE SPECIFICITY: KIDNEY, INTESTINAL BRUSH BORDER, AND SALIVARY DUCTS.
CC					CC -1- SIMILARITY: BELONGS TO PEPTIDASE FAMILY M12A (ZINC METALLOPROTEASE); ALSO KNOWN AS THE ASTACIN SUBFAMILY.
CC					CC -1- SIMILARITY: CONTAINS 1 MAM DOMAIN.
Db	430 CRYCLRLKCFRAGMKKEAVONERDRISTRSTFDG 464				RESULT 10 ID YB9K_YEAST STANDARD; PRT; 320 AA.
AC	1:1:1 1: :1   :1   :: : ;       :				DT 01-OCT-1994 (Rel. 30, Created)
DT	01-OCT-1994 (Rel. 30, Last sequence update)				DT 01-OCT-1994 (Rel. 30, Last annotation update)
Qy	45 CWTCCRNSYRALDK-SLHVGTQCALTRCPQEG 78				DE HYPOTHETICAL PROTEIN IN SHM1-MRRL37 INTERGENIC REGION.
OS	Saccharomyces cerevisiae (Baker's yeast).				GN YBR265W OR YBR1734.
OC	Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;				CC



DT 01-JAN-1988 (Rel. 06, Created)  
 DT 01-DEC-1992 (Rel. 24, Last sequence update)  
 DT 15-JUL-1999 (Rel. 38, Last annotation update)  
 DE T-CELL SURFACE GLYCOPROTEIN CD1A PRECURSOR (CD1A ANTIGEN) (T-CELL SURFACE ANTIGEN T6/LEU-6) (HTAL THYMOCYTE ANTIGEN).  
 GN CD1A.  
 OS Homo sapiens (Human).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 [1] RN SEQUENCE FROM N.A.  
 RX MEDLINE: 88097453.  
 RA ARUFFO A.; SEED B.;  
 RT "EXPRESSION OF cDNA clones encoding the thymocyte antigens CD1a, b, c demonstrates a hierarchy of exclusion in fibroblasts.";  
 RT J. Immunol. 143:1723-1730(1989).  
 [3] RN SEQUENCE OF 99-327 FROM N.A.  
 RP TISSUE=T-CELL;  
 RX MEDLINE: 87014824.  
 RA CALABI F.; MILSTEIN C.;  
 RT "A novel family of human major histocompatibility complex-related genes not mapping to chromosome 6.";  
 RL Nature 323:540-543(1986).  
 CC -1- SUBUNIT: ASSOCIATES NON-COVALENTLY WITH BETA-2-MICROGLOBULIN.  
 CC -1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.  
 CC -1- TISSUE SPECIFICITY: EXPRESSED ON CORTICAL THYMOCYTES, ON CERTAIN T-CELL LEUKEMIAS, AND IN VARIOUS OTHER TISSUES.  
 CC -1- SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN SUPERFAMILY.  
 CC -1- DATABASE: NAME-PROW; NOTE-CD guide CD1 entry;  
 CC WWW="http://www.ncbi.nlm.nih.gov/prov/cd/cd1.htm".

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DR EMBL; M28625; AAA51931.1; .  
 DR EMBL; M22167; AAA51932.1; .  
 DR EMBL; M22080; AAA51932.1; JOINED.  
 DR EMBL; M22163; AAA51932.1; JOINED.  
 DR EMBL; M22164; AAA51932.1; JOINED.  
 DR EMBL; M22165; AAA51932.1; JOINED.  
 DR EMBL; M22166; AAA51932.1; JOINED.  
 DR EMBL; X04450; CAA28049.1; .  
 DR PIR; A02242; HLHUCD.  
 DR PIR; A39957; A39957.  
 DR HSSP; P11609; 1CD1.  
 DR MIM; 188370; .  
 DR PFAM; PF00047; ig\_1; Glycoprotein; Signal; Transmembrane; Immunoglobulin domain; T-cell; Multigene family.  
 KW Multigene family.

SIGNAL 1 16  
 FT CHAIN 17 327  
 FT DOMAIN 17 108  
 FT DOMAIN 109 201  
 FT DOMAIN 202 294  
 FT TRANSMEM 300 321  
 FT DOMAIN 322 327  
 FT DISULFID 119 183  
 FT DISULFID 223 278

FT CARBOHYD 37 37  
 FT CARBOHYD 60 60  
 FT CARBOHYD 74 74  
 FT CARBOHYD 145 145  
 FT CONFLICT 30 30  
 FT CONFLICT 68 68  
 SQ SEQUENCE 327 AA; 37172 MW; 613EF65 CRC32;

Query Match 9.4%; Score 84; DB:1; Length 327;  
 Best Local Similarity 48.1%; Pred. No. 1.09e+00;  
 Matches 13; Conservative 7; Mismatches 4; Indels 3; Gaps 3;

DB 299 SVGFILLAVIVPLLLIGLWF-RKR 324  
 QY 27 AAGIGIIVLVILGVLLIG-C-WCRRR 51

RESULT 14  
 ID Y472\_RICPR STANDARD PRT; 334 AA.  
 AC Q9ZD72;  
 DT 15-DEC-1999 (Rel. 39, Created)  
 DT 15-DEC-1999 (Rel. 39, Last sequence update)  
 J. Immunol. 143:1723-1730(1989).  
 DE HYPOTHETICAL PROTEIN RP472.  
 GN RP472.  
 OS Rickettsia prowazekii.  
 OC Bacteria; Proteobacteria; alpha subdivision; Rickettsiales;  
 OC Rickettsiaceae; Rickettsiae; Rickettsia.  
 RN 11  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=MADRID E;  
 RX MEDLINE: 99039499.  
 RA ANDERSSON S.G.E.; ZOMORDIPOUR A.; ANDERSSON J.O.;  
 RA SICHERITZ-PONTEN T.; ALSMARK U.C.M.; PODOWSKI R.M.; NAESLUND A.K.;  
 RA ERIKSSON A.-S.; WINKLER H.H.; KURLEND C.G.;  
 RT "THE GENOME SEQUENCE OF Rickettsia prowazekii AND THE ORIGIN OF MITOCHONDRIA";  
 RT Nature 396:133-140(1998).  
 CC DR AU235271; CAA14927.1; .  
 CC KW HYPOTHETICAL protein; Transmembrane.  
 CC FT TRANSMEM 1 21 POTENTIAL.  
 CC FT TRANSMEM 46 60 96F50415 CRC32;  
 CC SQ SEQUENCE 334 AA; 39111 MW; 96F50415 CRC32;

Query Match 9.4%; Score 84; DB:1; Length 334;  
 Best Local Similarity 64.3%; Pred. No. 1.09e+00;  
 Matches 9; Conservative 5; Mismatches 0; Indels 0; Gaps 0;

DB 42 FLSVIGLGLLLYSC 55  
 QY 32 ILTVILGVLLIGC 45

RESULT 15  
 ID C1080\_MOUSE STANDARD PRT; 306 AA.  
 AC Q00609;  
 DT 01-FEB-1994 (Rel. 28, Created)  
 DT 01-FEB-1994 (Rel. 28, Last sequence update)  
 DT 01-FEB-1996 (Rel. 33, Last annotation update)  
 DE T LYMPHOCYTE ACTIVATION ANTIGEN CD80 PRECURSOR (ACTIVATION B7-1 DE ANTIGEN) (B7).  
 GN C1080 OR B7.  
 OS Mus musculus (Mouse).  
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus .  
 RN [1] SEQUENCE FROM N.A.  
 RP TISSUE-B CELL;  
 RC MEDLINE; 91311422.  
 RX GRAY G.S.; FREEMAN G.J.; GIMMI C.D.; LOMBARD D.B.; ZHOU L.J.;  
 RA WHITE M.; FINGEROTH J.D.; GRIBBEN J.G.; NADLER L.M.;  
 RT "Structure, expression, and T cell costimulatory activity of the  
 murine homologue of the human B lymphocyte activation antigen B7.";  
 RT J. Exp. Med. 174:625-631(1991).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC TISSUE-B-CELL;  
 RX MEDLINE; 93307789.  
 RA SELVIKUMAR A.; WHITE P.C.; DUPONT B.;  
 RT Genomic organization of the mouse B-lymphocyte activation antigen  
 RL Immunogenetics 38:292-295(1993).  
 CC -1- FUNCTION: INVOLVED IN THE COSTIMULATORY SIGNAL ESSENTIAL FOR T  
 CC LYMPHOCTYES ACTIVATION. T CELL PROLIFERATION AND CYTOKINE  
 CC PRODUCTION IS INDUCED BY THE BINDING OF CD28 OR CTLA-4 TO THIS  
 CC RECEPTOR.  
 CC -1- SUBCELLULAR LOCATION: TYPE I MEMBRANE PROTEIN.  
 CC -1- TISSUE SPECIFICITY: EXPRESSED ON ACTIVATED B CELLS, GAMMA  
 CC INTERFERON STIMULATED MONOCYTES AND NONCIRCULATING B-CELL  
 CC MALIGNANCIES.  
 CC -1- DEVELOPMENTAL STAGE: EXPRESSED BETWEEN 4 AND 12 HOURS POST-  
 CC ACTIVATION. PROTEIN WAS DETECTED AT CELL SURFACE AT 24 HOURS AND  
 CC IT'S EXPRESSION WAS MAXIMAL FROM 48 TO 72 HOURS POST-ACTIVATION.  
 CC -1- SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN SUPERFAMILY. CONTAINS  
 CC ONE C2-LIKE AND ONE V-LIKE DOMAINS.  
 CC -----  
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration  
 CC between the Swiss Institute of Bioinformatics and the EMBL outstation -  
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 CC or send an email to [license@isb-sib.ch](mailto:license@isb-sib.ch)).  
 CC -----  
 DR X60958; CAA43291.1;  
 DR EMBL; L11589; AAA37240.1; ALT SEQ.  
 DR EMBL; L12585; AAA37240.1; JOINED.  
 DR EMBL; L12586; AAA37240.1; JOINED.  
 DR EMBL; L12587; AAA37240.1; JOINED.  
 DR EMBL; L12588; AAA37240.1; JOINED.  
 DR PIR; S17291; S17291.  
 DR MGI; MGI:101775; CD80.  
 DR PFAM; PF00047; 19; 2.  
 KW Immuno-globulin domain; T-cell; Glycoprotein; Signal; Transmembrane;  
 KW Receptor.  
 FT SIGNAL 1 37  
 FT CHAIN 38 306 T LYMPHOCYTE ACTIVATION ANTIGEN CD80 .  
 FT DOMAIN 38 246 EXTRACELLULAR (POTENTIAL).  
 FT TRANSMEM 247 268 POTENTIAL.  
 FT DOMAIN 269 306 CYTOPLASMIC (POTENTIAL).  
 FT DOMAIN 47 126 IG-LIKE V-TYPE DOMAIN.  
 FT DOMAIN 158 226 IG-LIKE C2-TYPE DOMAIN.  
 FT DOMAIN 227 246 IG-HINGE LIKE (POTENTIAL).  
 FT DISULFID 54 119 POTENTIAL.  
 FT DISULFID 165 219 POTENTIAL.  
 FT CARBOHYD 93 93 POTENTIAL.  
 FT CARBOHYD 99 99 POTENTIAL.  
 FT CARBOHYD 149 149 POTENTIAL.  
 FT CARBOHYD 189 189 POTENTIAL.  
 FT CARBOHYD 210 210 POTENTIAL.  
 FT CARBOHYD 214 214 POTENTIAL.  
 SQ SEQUENCE 306 AA; 34589 MW; 86FDD183 CRC32;  
 Query Match 9.3%; Score 83; DB 1; Length 306;  
 Best Local Similarity 27.6%; Pre. No. 1.5e+00;  
 Matches 8; Conservative 14; Mismatches 6; Indels 1; Gaps 1;

Secreted protein; human; cell proliferation; cytokine activity;  
 KW tissue growth; cellular differentiation; regeneration; activity;  
 KW inhibin; chemotactic; haemostatic; thrombolytic; tumour inhibition;  
 KW anti-inflammatory activity; biomarker.  
 OS Homo sapiens.  
 PN WO925959-A2.  
 PD 18-JUN-1998.  
 PF 11-DEC-1997; 022787.  
 PR 11-DEC-1996; US-032757.  
 PA (CHIR) CHIRON CORP.  
 PI Escobedo J., Garcia P., Hu Q., Rothakota S., Williams LT;  
 DR WPI: 98-348453/30.  
 DB N-PSDB: V43602.  
 PT Secreted human polypeptides - having cytokine, cell proliferation or  
 PT differentiation, activin or inhibin, tumour inhibition or  
 PT anti-inflammatory activities  
 PS Claim 1; Pages 49-50; 78pp; English.  
 CC This represents a human secreted protein. The specification provides  
 CC secreted protein sequences (WO3681 to WO3699) encoded by the nucleic  
 CC acid sequences shown in V43601 to V43619. The invention provides a  
 CC method of identifying a secreted polypeptide which is modified by rough  
 CC microsomes. The secreted proteins can be used in assays to determine  
 CC biological activities, such as cytokine, cell proliferation, or cellular  
 CC differentiation activities, tissue growth or regeneration, activin or  
 CC inhibin activity, chemotactic or chemokinetic activity, haemostatic or  
 CC thrombolytic activity, receptor/ligand activity, tumour inhibition, or  
 CC anti-inflammatory activity. The proteins can also be used as biomarkers,  
 CC to identify tissues or cell types which express the proteins, or a stage-  
 CC or disease-specific alteration in protein expression. They can be used  
 CC in protein interaction assays, to identify ligands or binding proteins.  
 CC Compounds which affect the biological activities of the secreted proteins  
 CC or their ability to interact with specific ligands can be identified  
 CC using the proteins in screening assays. The proteins and antibodies that  
 CC bind specifically to the protein can also be used to design diagnostic  
 CC tests and therapeutic compositions for diseases which may be associated  
 CC with altered expression of these proteins. Fusion proteins comprising,  
 CC e.g. signal sequences or transmembrane domains of the proteins can be  
 CC used to target other protein domains to cellular membrane or they can  
 CC be secreted extracellularly.  
 Sequence 291 AA;

Query Match 6 standard; protein: 380 AA.  
 Best Local Similarity 10.6%; Pred. No. 5.23e+00;  
 Matches 17; Conservative 7; Mismatches 4; Indels 4; Gaps 4;  
 Db 257 ALAVAVLKTVIGLGLCLL-LWW-RRRKGSRA 286  
 Qy 27 AAGIGL-TVIGLV-LIGCWICRRRNGYRA 56

RESULT 6 R05433 standard; protein: 380 AA.  
 ID R05433.  
 AC R05433.  
 DT 30-JUL-1990 (first entry)  
 DE CPA-P2 Hybrid plasminogen activator.  
 KW Plasminogen activator; fibrin; urokinase; thromboembolic  
 KW disease; ds.  
 OS Synthetic.  
 FH Location/Qualifiers  
 ECdomain 1..23  
 FT domain 1..102  
 FT domain 103..116  
 FT domain 117..380  
 FT domain 1..116  
 FT domain 117..380  
 PN WO9001332-A.  
 PD 22-FEB-1990.  
 PF 10-AUG-1988; 02771.  
 PR 10-AUG-1988; WO-022771.  
 PA (Cetus) Cetus Corp.  
 PI Halluin AP;

DR WPI: 90-083374/11.  
 DR N-PSDB: Q0201.  
 PT Compsn. config. plasminogen activator conjugated to heparin component -  
 PT used for treatment of thromboembolic disease, with longer half life and improved targeting.  
 PT Disclosure; P; English.  
 PS Gene encodes hybrid plasminogen activator (PA) comprising Kringle 1, an  
 CC urokinase linker, and an urokinase protease domain wherin glycine residue  
 CC at position 158 is replaced with a lysine.  
 CC The compound is used to treat thromboembolic disease esp. with myocardial  
 CC infarction, has a longer half-life than free PA and targets the heparin  
 CC site of thrombus or embolism reducing the risk of reocclusion.  
 SQ sequence 380 AA;

Query Match 10.1%; Score 90; DB 1; Length 380;  
 Best Local Similarity 30.9%; Pred. No. 1.1e+01;  
 Matches 21; Conservative 15; Mismatches 24; Indels 8; Gaps 7;

Db 7 LAALLLILLLPGCGWASSEKTKGDKKNYRTMSKPKN-GTTCQKWSSTSHPRPRESPATHPS 65  
 Qy 33 IIVLGVLVLLGCV-Y-CRRRG-TRALMDLSLVGTQCAL-TRRQEQ-FDH-RDSB 85

RESULT 7 W00407 standard; Protein: 226 AA.  
 ID W00407.  
 AC W00407.  
 DT 13-JAN-1999 (first entry)  
 DE A secreted protein encoded by clone di39\_9.  
 KW secreted protein; immune stimulating; suppressing;  
 KW haematopoiesis regulating activity; tissue growth activity; activin;  
 KW inhibin activity; chemotactic; chemokinetic activity; haemostatic;  
 KW thrombolytic activity; anti-inflammatory activity; cadherin;  
 KW tumour invasion suppressor activity; tumour inhibition activity.  
 OS Homo sapiens.

PD 08-OCT-1998.  
 PF 27-MAR-1998; U06176.  
 PR 28-MAR-1997; US-823330.  
 PA (GEMY ) GENETICS INST INC.

PI Agostino MJ, Jacobs K, Lavallie ER, MCCOY JM, Merberg D,  
 PI Racine LA, Spaulding V, Treacy M,  
 DR WFI: 98-54203/46.  
 DR N-PSDB: V63191.

PT New isolated polynucleotide(s) and secreted proteins - are obtained  
 PT from human cDNA libraries prepared from adult testes, foetal brain,  
 PT adult brain, adult blood and placenta.  
 PS Claim 19; Pages 76-77; 124pp; English.  
 CC The present sequence represents a secreted protein. The nucleic acid  
 CC sequence is isolated from a human adult testes cDNA library using  
 CC probe V63202. The polypeptide may have biological activities such has  
 CC e.g. nutritional activity, immune stimulating or suppressing activity,  
 CC haematopoiesis regulating activity, tissue growth activity;  
 CC activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic  
 CC and thrombolytic activity, receptor/ligand activity, anti-inflammatory  
 CC activity, cadherin/tumour invasion suppressor activity, tumour inhibition  
 SQ Sequence 226 AA;

Query Match 10.0%; Score 89; DB 1; Length 226;  
 Best Local Similarity 53.1%; Pred. No. 1.34e+01;  
 Matches 17; Conservative 7; Mismatches 3; Indels 5; Gaps 4;

Db 193 ALAVAVLKTVIGLGLCLL-LWW-RRRKGSRA 221  
 Qy 27 AAGIGL-TVIGLV-LIGCWICRRRNGYRA 56

W20924 standard; protein; 141 AA.  
 W20924;  
 AC ID  
 AC 21-JUL-1997 (first entry)  
 DE H. pylori cell envelope protein, 16ae10505orf13.  
 KW cytoplasmic; vaccine; prevention; treatment; infection; identification;  
 binding compound; bacterium; life cycle; activator; bacteria; inhibitor;  
 duodenal ulcer disease; chronic gastritis; diagnosis; envelope.  
 KW Helicobacter pylori.  
 OS Helicobacter pylori.  
 PN W09640893-A1.  
 PN T-DEC-1996.  
 PD 06 JUN 1996; U09122.  
 PF 07 JUN 1995; US 4,870,322.  
 PR 07 JUN 1995; US 4,870,322.  
 PR 01 APR 1996; US 6,304,055.  
 PR (ASTR ) ASTRA AB.  
 PA Berglindh OT; Smith D, Mellgaard BL;  
 PI DR WPI; 97-052306,05.  
 DR N-PSDB; T81177.  
 DR Helicobacter pylori nucleic acid sequences and related  
 polypeptide(s) - useful for vaccines to treat or prevent H. pylori  
 infection, and to detect Helicobacter  
 PT infection. and to detect Helicobacter  
 PS Claim 56: Page 1322; 1481pp; English.  
 CC The present sequence is a H. pylori cell envelope protein.  
 CC The protein may be used in a vaccine to prevent or treat H. pylori  
 CC infection or to identify H. pylori polypeptide binding compounds,  
 CC useful as potential H. pylori life cycle activators or inhibitors.  
 CC The genomic sequence of H. pylori (ATCC 55679) was determined from  
 CC overlapping contigs generated by mechanically shearing the bacterial  
 CC DNA. The sequences were analysed for ORF of at least 180 nucleotides,  
 CC and the predicted coding regions defined by computer evaluation. To  
 CC identify likely H. pylori antigens for vaccine development, the amino  
 CC acid sequences predicted from various ORF were analysed for significant  
 CC homology to other known or exported membrane proteins. Having identified  
 CC and determined the sequences of interest, particular regions can be  
 CC isolated from H. pylori by PCR amplification for recombinant polypeptide  
 CC

CC 8 nucleotides from the nucleic acid sequences. The vaccines are useful  
CC for treating or reducing the risk of *H. pylori* infections, and the  
CC probes can be used diagnostically for detecting the presence of  
CC *Helicobacter* in a sample. The products are also of use in screening  
CC for compounds having the ability to interfere with the *H. pylori* life  
CC cycle or to inhibit *H. pylori* infection.  
CC 169 AA;  
CC Sequence:

Query	Match	Score	Length
Db	1.04	AEGLGRITIMIGLIVLGLW	123
Qv	27	AAGIGTIVLGVVLLGGW	46
Best	9.9%	Score 88;	DB 1;
Local	55.0%	Pred. No. 1.	6.1e+01;
Similarity		Matches	Lengt
Matches	11;	Conservative	169;
		Mismatches	
		Indels	
		0;	Gaps

N-Prbus; 1081/7). Helicobacter pylori nucleic acid sequences and related polypeptide(s) - useful for vaccines to treat or prevent H. pylori infection, and to detect Helicobacter pylori. Page 1322; 1481PP; English. Claim 56; Page 1322; 1481PP; English.

The present sequence is a H. pylori cell envelope protein. The protein may be used in a vaccine to prevent or treat H. pylori infection or to identify H. pylori polypeptide binding compounds, useful as potential H. pylori life cycle titrators or inhibitors. The genomic sequence of H. pylori (ATCC 55679) was determined from overlapping contigs generated by mechanically shearing the bacterial DNA. The sequences were analysed for ORF of at least 180 nucleotides, and the predicted coding regions defined by computer evaluation. To identify likely H. pylori antigens for vaccine development, the amino acid sequences predicted from various ORF were analysed for significant homology to other known or exported membrane proteins. Having identified and determined the sequences of interest, particular regions can be isolated from H. pylori by PCR amplification for recombinant polypeptide

PT Helicobacter pylori nucleic acids and proteins - used to develop  
PT products for the detection, prevention and treatment of *H. pylori*  
PT infections  
PS Claims 27, 31; Page 226-227; 27pp; English.  
CC Recombinant or substantially pure preparations of *H. pylori* polypeptides  
CC are disclosed, together with the nucleic acids encoding them. In all,  
CC 73 ORFs are shown. The proteins are variously cell envelope proteins,  
CC secreted proteins or other cellular proteins. Vaccines containing the  
CC nucleic acids or proteins are claimed, as are probes containing at least  
CC 8 nucleotides from the nucleic acid sequences. The vaccines are useful  
CC for treating or reducing the risk of *H. pylori* infections, and the  
CC probes can be used diagnostically for detecting the presence of  
CC *Helicobacter* in a sample. The products are also of use in screening  
CC for compounds having the ability to interfere with the *H. pylori* life  
CC cycle or to inhibit *H. pylori* infection.  
SO Sequence 215 AA:  
SQ

Query	Match	Score 88;	DB 1;	Length 215;
Best Local	Similarity	9.9%	Pred. No.	1.6e+01;
Matches	11;	Conservative	Mismatches	5;
			Indels	0;
			Gaps	
Db	104	AEGILGITIMILGLIVLGLW	123	
Ov	27	AAGCTTGTTCATGTTCTGG	46	

PA (ASTRA AB.)  
PI Alm RA, Smith D;  
WPI; 98-271811/24.  
N-PSDB: X30474.  
DR Helicobacter pylori nucleic acids and proteins - used to develop  
PT products for the detection, prevention and treatment of *H. pylori*  
PT infections  
Pr Claims 27, 31; Page 217; 279pp; English.  
PS Recombinant or substantially pure preparations of *H. pylori* polypeptides  
CC are disclosed, together with the nucleic acids encoding them. In all,  
CC 73 ORPs are shown. The proteins are variously cell envelope proteins,  
CC secreted proteins or other cellular proteins. Vaccines containing the  
CC nucleic acids or proteins are claimed, as are probes containing at least  
CC one of the nucleic acids.

/note= "antigenic peptide sequence recognised by CTL"

WT W09729195-A2.  
PN 14-AUG-1997; U02186;  
PD 06-FEB-1997; U02186;  
PF 04-OCT-1996; US-725736.  
PR 09-FEB-1996; US-599602.  
PA (USSN ) US DEPT HEALTH & HUMAN SERVICES.  
PI Rosenberg SA, Wang R;  
DR WPI; 97-15349/38.  
DR N-PSDB; T91957.  
PT Cancer antigen peptide(s) derived from the tyrosinase-related protein 1 or 2 - useful for detecting, preventing or treating a cancer in a mammal, especially melanoma  
PT Claim 12; Pages 81-83; 11pp; English.  
CC The present sequence represents the novel tyrosinase related protein 2 (TRP-2). This protein contains tumour antigens recognised by tumour infiltrating lymphocyte (TIL) 586. Novel cancer peptides have also been identified in TRP-1. The peptides are recognised by a major histocompatibility complex (MHC) class I T-lymphocyte. The nucleic acids encoding the cancer peptides or TRP-2 can be used to detect a cancer or pre-cancer in a mammal, especially by detecting the presence of the alternative ORF 3 of the TRP-1 gene or the sequence encoding the novel tumour antigen TRP-2. Vectors and recombinant viruses containing antigen peptide encoding nucleic acids, antibodies raised against the peptides, or the peptides themselves can be used to prevent or treat a cancer in a mammal, especially a melanoma.  
SQ Sequence 519 AA;

Query Match 9.8%; Score 87; DB 1; Length 519;  
Best Local Similarity 35.3%; Pred. No. 1.94e+01;  
Matches 12; Conservative 11; Mismatches 10; Indels 1; Gaps 1;

Db 477 MGTIVAVLQGLFLVLLAFQLQYRRLKRYTIPMETHL 510  
QY 30 IGTITVLIGLVLLIGCWCRR-RNGYRALMDKS 62

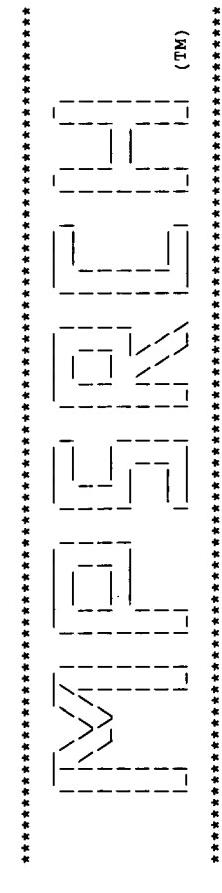
RESULT 12  
ID W20802 standard; protein; 296 AA.  
AC W20802;  
DT 16-JUL-1997 (first entry)  
DE H. pylori inner membrane protein, Ogap11406orf5.  
KW Cyttoplasmic; vaccine; prevention; treatment; infection; binding compound; bacterium; life cycle; activator; bacteria; inhibitor; diabulon ulcer disease; chronic gastritis; diagnosis; envelope.  
KW Helicobacter pylori.  
OS W09640893-A1.  
PN 19-DEC-1996.  
PD 06-JUN-1996; U09122.  
PR 07-JUN-1995; US-487032.  
PR 01-APR-1996; US-630405.  
PA (ASPR ) ASTRA AB.  
PI Berglindh OT, Smith D, Mellgaard BL;  
DR N-PSDB; T68055.  
PT Helicobacter pylori nucleic acid sequences and related polypeptide(s) - useful for vaccines to treat or prevent H. pylori infection, and to detect Helicobacter

PS 56; Page 1209; 1401pp; English.  
CC The present sequence is a H. pylori inner membrane protein.  
CC The protein may be used in a vaccine to prevent or treat H. pylori infection to identify H. pylori polypeptide binding compounds, useful as potential H. pylori life cycle activators or inhibitors.  
CC The genomic sequence of H. pylori (ATCC 55673) was determined from overlapping contigs generated by mechanically shearing the bacterial DNA. The sequences were analysed for ORF of at least 180 nucleotides, and the predicted coding regions defined by computer evaluation. To identify likely H. pylori antigens for vaccine development, the amino acid sequences predicted from various ORF were analysed for significant homology to other known or exported membrane proteins. Having identified and determined the sequences of interest, particular regions can be isolated from H. pylori by PCR amplification for recombinant polypeptide production, e.g. in E. coli hosts.

SQ	Sequence	296 AA;
Query	Match	9.4%; Best Local Similarity 29.7%; Matches 11; Conservative 14; Mismatches 11; Indels 1; Gaps 1;
Db	210 FFGNLKTGNNOISVFEDLNAREGVLSVTLALIIG 246 QY 9 ITCYPKKGHGSYTTAE-EAAGIGLTVLGVLILLG 44	
RESULT 13		
ID	R82900	standard; Protein; 200 AA.
AC	R82900;	
DT	07-MAY-1996 (first entry)	
DE	Mouse B7-1 (IgV-like domain deleted).	
KW	T-cell co-stimulatory molecule; B7-1; T-lymphocyte; CD28; CTLA4; receptor; immunoglobulin.	
OS	Mus musculus.	
Location/Qualifiers		
FH	Key	
FT	peptide	1..37
FT	domain	169..200
FT	label= "Cytoplasmic_domain	
FT	note= "cytoplasmic_domain is encoded by exon 5 of the B7-1 gene"	
FT	FT	
PN	W09523859-A2.	
PD	08-SEP-1995;	
PF	02-MAR-1995; U02576.	
PR	02-MAR-1994; US-205697.	
PA	(BCHM ) BRIGHAM & WOMENS HOSPITAL.	
PA	(DAND ) DANA FARBER CANCER INST.	
PI	Borriello F, Freeman GJ, Nadler LM, Sharpe AH;	
DR	95-320574/41.	
PT	Novel T cell co-stimulatory molecules - corresponding to naturally occurring alternatively spliced forms of T cells co-stimulatory molecules or variants	
PT	PS Disclosure; Page 55-56; 11pp; English.	
CC	A naturally occurring form of mouse T-cell costimulatory molecule B7-1 (R82900) has the signal peptide directly linked to the IgV-like domain, i.e. the IgV-like domain is deleted. It is encoded by exons 1, 3, 4 and 5 (see T01047) of the B7-1 gene. An alternatively spliced form of IgV-deleted B7-1 (R82901) is encoded by exons 1, 3, 4 and 6. T-cell costimulatory molecules can be produced in which the IgV-like domain is deleted.	
SQ	Sequence 200 AA;	
Query	Match	9.3%; Best Local Similarity 27.6%; Matches 8; Conservative 14; Mismatches 6; Indels 1; Gaps 1;
Db	146 GAGFGAVITVVVIVVVIKCFCKHRSPFR 173 QY 27 AAGIGITLVIGLVLLIGCWCRRNGYR 55	
RESULT 14		
ID	R82902 standard; Protein; 212 AA.	
AC	R82902;	
DT	07-MAY-1996 (first entry)	
DE	Mouse B7-1 IgV-like isoform.	
KW	T-cell co-stimulatory molecule; B7-1; T-lymphocyte; CD28; CTLA4; receptor; immunoglobulin; interleukin-2.	
OS	Mus musculus.	
Location/Qualifiers		
FH	Key	1..37
FT	peptide	1..212
FT	label= "Cytoplasmic_domain	
FT	note= "cytoplasmic_domain is encoded by exon 5 of the B7-1 gene"	
PN	W09523859-A2.	

PD 08-SEP-1995.  
 PF 02-MAR-1995; UD2576  
 PF 02-MAR-1994; US-205697.  
 PA (BGHM ) BRIGHAM & WOMENS HOSPITAL.  
 PA \*(DAND ) DANA FARBER CANCER INST.  
 PI 'Borriello F, Freeman GJ, Nadler LM, Sharpe AH;  
 WPI 95-320474/41.  
 DR N-PSDB: T01049.  
 DR Novel T cell co-stimulatory molecules - corresponding to naturally  
 occurring alternatively spliced forms of T cells co-stimulatory  
 molecules or variants  
 Disclosure; Page 91-92; 111pp; English.  
 A naturally occurring form of mouse T-cell costimulatory molecule  
 B7-1 (R82902) has the IgY-like domain directly linked to the  
 transmembrane domain, i.e. the IgC-like domain is deleted. It is  
 encoded by exons 1, 2, 4 and 5 of the B7-1 (T01049) gene.  
 This IgY-like isoform of B701 was expressed in CHO cells. It  
 triggered a costimulatory signal in T-cells, causing stimulation  
 of interleukin-2 prodn.  
 Sequence 212 AA;  
 CC SQ

RESULT	15	ID	R67990 standard; Protein; 306 AA.
		AC	R67990;
		DT	21-AUG-1995 (first entry)
		DE	Murine B lymphocyte antigen B7 (mbB7).
		KW	B lymphocyte activation antigen; B7-1; Ig superfamily; CD28; transmembrane protein.
		OS	Mus musculus.
		FH	Key
		FT	protein
		FT	
		FT	label= signal sequence
		FT	/note= "hydrophobic"
		domain	
		FT	38 . .247
		FT	label= extracellular
		FT	/note= "6"
		domain	
		FT	248 . .272
		FT	label= transmembrane
		domain	
		FT	273 . .306
		FT	label= intracellular (cytoplasmic)
		domain	
		FT	38 . .142
		FT	label= Ig V-set domain
		domain	
		FT	143 . .236
		FT	label= Ig C-set domain
		FT	misc_difference 1 . .306



Release 3.1A John F. Collins, Biocomputing Research Unit.

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Mpsrch\_pp protein - protein database search, using Smith-Waterman algorithm

Run on: Fri May 5 22:01:50 2000; MasPar time 7.16 Seconds

Tabular output not generated.  
87.109 Million cell updates/sec

Title: >US-09-267-439-4

Description: (1-9) from US09267439 .pep

Perfect Score: 56

Sequence: 1 AAAGIILTV 9

Scoring table: PAM 150  
Gap 15

Searched: 225878 seqs, 69334122 residues

Post-processing: Minimum Match 0<sup>8</sup>  
Listing first 45 summaries

Database: sptrembl12

1:sp\_archea 2:sp\_bacteria 3:sp\_fungi 4:sp\_human  
5:sp\_invertebrate 6:sp\_mammal 7:sp\_mhc 8:sp\_organelle  
9:sp\_phage 10:sp\_plant 11:sp\_rabbit 12:sp\_unclassified  
13:sp\_vertebrate 14:sp\_yivirus

Statistics: Mean 21.729; Variance 25.002; scale 0.869

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query	Match	Length	DB	ID	Description	Pred. No.
1	49	87.5	766 10	023161	RECEPTOR KINASE-LIKE P	3.6e+00		
2	48	83.7	808 10	Q9ZT37	PUTATIVE GLUTAMATE REC	6.11e+00		
3	47	83.9	165 2	P71810	HYPOTHETICAL 18.2 KD P	1.01e+01		
4	47	83.9	250 2	Q31597	KJBA PROTEIN	1.01e+01		
5	47	83.9	478 14	Q87090	GLYCOPROTEIN GLII.	1.01e+01		
6	47	83.9	479 14	Q87089	GLYCOPROTEIN GLII.	1.01e+01		
7	47	83.9	479 14	Q87091	GLYCOPROTEIN GLII.	1.01e+01		
8	46	82.1	848 5	Q18139	T26H2.7 PROTEIN	1.67e+01		
9	45	80.4	339 1	Q30640	METHYLCOBAMIDE-COM MET	2.73e+01		
10	45	80.4	339 1	Q48928	METHYLCOBAMIDE-COM MET	2.73e+01		
11	45	80.4	339 1	Q48950	METHYLCOBALAMIN: COENZ	2.73e+01		
12	45	80.4	370 8	Q48172	CYTOKRONE B.	2.73e+01		
13	45	80.4	420 11	Q63276	KAN-1.	2.73e+01		
14	45	80.4	420 11	Q08833	BILE ACID COA: AMINO A	2.73e+01		
15	45	80.4	980 5	Q17592	SIMILARITY TO INSULIN-XYLANSIE	2.73e+01		
16	45	80.4	1347 2	Q30426	HYPOTHETICAL 10.3 KD P	4.43e+01		
17	44	78.6	98 2	Q928X3	ABC TRANSPORTER, ATP-B	4.43e+01		
18	44	78.6	620 1	Q29198	ACETYLCHOLINE RECEPTOR	7.14e+01		
19	43	76.8	91 11	Q54712	F13F21.20 PROTEIN.	7.14e+01		
20	43	76.8	190 10	Q9XIA3				

#### ALIGNMENTS

```
RESULT 1 PRELIMINARY;
ID 023161;
AC Q33161;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-MAY-1999 (TREMBLrel. 10, Last sequence update)
DE RECEPTOR KINASE-LIKE PROTEIN (EC 2.7.1.1.)
GN C/A10.110.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophytina; Embryophyta; Tracheophyta;
OC euphyllophytes; Spermato phyta; Magnoliophyta; eudicotyledons;
OC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;
OC Arabidopsis.
RN [1]
RP SEQUENCE FROM N.A.
RA BRIAN M., TERRYN N., VOS P., HELIJNEN L., NEWES H.W., SCHUELLER C.,
RA CHALWATZIS N.; Submitted (DEC-1998) to the EMBL/GenBank/DDBJ databases.
RL DR; 29707; CAB16774.1; DR; 25436; Arath; 3435; 25486.
DR MENDEL; PF0050; LRR; 4; DR; PFAM; PF0050; Pfam; 1.
DR PRINTS; /PRO0019; LEURCHRPT.
SEQUENCE 766 AA; 83775 MW; C4BD7115 CRC32;
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Query/Match Similarity 87.5%; Best/Local Similarity 87.5%; Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
Db' 337 AGIGILAY 344
| | | | | |
Qy 2 AGIGILTV 9
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RESULT 2 PRELIMINARY;
ID Q9ZT37;
AC Q9ZT37;
DT 01-MAY-1999 (TREMBLrel. 10, Created)
DT 01-MAY-1999 (TREMBLrel. 10, Last sequence update)
DE PUTATIVE GLUTAMATE RECEPTOR.
GN GIRL.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophytina; Embryophyta; Tracheophyta;
```

OC euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;  
 OC core eudicots; Rosidae; eurosids-II; Brassicales; Brassicaceae;  
 OC Aroidiopsis.  
 [1] RN SEQUENCE FROM N.A.  
 RX MEDLINE; 99039497.  
 RA LAM H., CHIU J., HSIEH M.H., METSEL L., OLIVEIRA I.C., SHIN M.,  
 RA CORUZZI G.;  
 RT "Glutamate-receptor genes in plants.";  
 RL Nature; 396:125-126(1998).  
 DR EMBL; AF079998; AAC09173; 1; -.  
 KW Receptor.  
 SQ SEQUENCE 808 AA; 90518 MW; C3554B89 CRC32;

Query Match 85.7%; Score 48; DB 10; Length 80;  
 Best Local Similarity 100.0%; Pred. No. 6.11e+00;  
 Matches 7; Conservative 0; Mismatches+0; Indels 0; Gaps 0;  
 RT

Db 519 GIGILTV 525  
 .3 GIGILTV 9

QC

RESULT 3 PRELIMINARY; PRT; 165 AA.  
 ID P71810; AC P71810; DT 01-FEB-1997 (TREMBLrel. 02, Created)  
 DT 01-NOV-1998 (TREMBLrel. 08, Last sequence update)  
 DE HYPOTHETICAL 18.2 KD PROTEIN.  
 GN MTCTB12.16.  
 OS Mycobacterium tuberculosis.  
 OC Bacteria; Firmicutes; Actinobacteria; Actinomycetidae;  
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.  
 [1] RN SEQUENCE FROM N.A.  
 RP STRAIN=H37RV;  
 RC MCLEAN J., HARRIS D.;  
 RA Submitted (OCT-1996) to the EMBL/GenBank/DBJ databases.  
 RL [2] SEQUENCE FROM N.A.  
 RP STRAIN=H37RV;  
 RA RAJANDREAM M.A.;  
 RL Submitted (OCT-1996) to the EMBL/GenBank/DBJ databases.  
 RN [3] SEQUENCE FROM N.A.  
 RP STRAIN=H37RV;  
 RX MEDLINE; 96181548.  
 RA PHILIP W.J., POULET S., EIGELMEIER K., PASCOPELLA L.,  
 RA BALASUBRAMANIAN V., HEYM B., BERGH S., BLOOM B.R., JACOBS W.R. JR.,  
 RA COLE S.T.;  
 RT "An integrated map of the genome of the tubercle bacillus,  
 leprae,";  
 RT Proc. Natl. Acad. Sci. U.S.A. 93:3132-3137(1996).  
 DR EMBL; 281011; CAB02643; 1; -.  
 KW Hypothetical protein.  
 SQ SEQUENCE 165 AA; 18189 MW; BFB84C79 CRC32;

Query Match 83.9%; Score 47; DB 2; Length 165;  
 Best Local Similarity 75.0%; Pred. No. 1.01e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

RESULT 4 PRELIMINARY; PRT; 250 AA.  
 ID 031597; AC 031597; DT 01-JAN-1998 (TREMBLrel. 05, Created)  
 DR 01-JAN-1998 (TREMBLrel. 05, Last sequence update)  
 SQ 2 AGIGILTV 135  
 2 AGIGILTV 9

RESULT 5 PRELIMINARY; PRT; 478 AA.  
 ID 037090; AC 037090; DT 01-NOV-1996 (TREMBLrel. 01, Created)  
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
 DE GLYCOPROTEIN GII.  
 OS Pseudorabies virus.  
 OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
 OC Alphaherpesvirinae; Varicellovirus.  
 RN 111  
 RP SEQUENCE FROM N.A.  
 RC SPAIN=INDIANA S;  
 RX MEDLINE; 96316347.  
 RA ISHIKAWA K., TSUTSUI M., TAGUCHI K., SAITO A., MURAMATSU M.;

DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)  
 DE YJBA PROTEIN.  
 GN YJBA.  
 OS Bacillus subtilis.  
 OC Bacteria; Firmicutes; Bacillus/Clostridium group;  
 OC Bacillus/Staphylococcus group; Bacillus.  
 [1] RN SEQUENCE FROM N.A.  
 RC STRAIN=168;  
 RX MEDLINE; 98044033.  
 RA KUNST F., OGASAWARA N., MOSZER I., ALBERTINI A.M., ALLONI G., BESSIRES P., BOLOTIN A., BORCHERT S., BRIGNEAU S.C., BRON S.,  
 RA AZEVEDO V., BERPERO M.G., BRAUN A., BRANS A., CAPOANO V., CALLOWELL B., CARTER N.M.,  
 RA BROUILLET S., BRUSCHI C.V., CONNERTON I.F., CUMMINGS N.-J., DANIEL R.A.,  
 RA CHOI S.K., CODANI J.J., CONNERTON I.F., DURRHOFF A., ERLICH S.D., EMMERSON P.T.,  
 RA DENIZOT F., DEVINE K.M., DERRINGTON J.J., FABRET C., FERRARI E., FOULGER D.,  
 RA ENTIAN K.D., FERRINGTON J.J., FUJITA M., FUJITA Y., FUMA S., GALIZZI A., GALLERON N.,  
 RA FRITZ C., FUJITA M., GOFFEUR A., GOLIGHTY E.J., GRANDI G.,  
 RA GUISSEPEI G., GUY B.J., HAGA K., HAIECH J., HARWOOD C.R., HENAUT A.,  
 RA HILBERT H., HOLSSAPPEL S., HOSONO S., HULLO M.F., ITAYA M., JONES L.,  
 RA JORIS B., KARAMATA D., KASAHARA Y., KLAER-BLANCHARD M., KLEIN C.,  
 RA KOBAYASHI Y., KOETTER P., KONINGSTEIN G., KROGH S., KUMANNO M.,  
 RA KURITA K., LAPIDUS A., LARDINOIS S., LAUBER J., LAZAREVIC V.,  
 RA LEE S.M., LEVINE A., LIU S., MAGUDA S., MAUEL C., MEDIGUE C.,  
 RA MEDINA N., MELLADO R.P., MIZUNO M., MOESTI D., NAKAI S., NOBACK M.,  
 RA NOONE D., O'REILLY M., OGAWA K., OGIVARA A., OODEGA B., PARK S.H.,  
 RA PARRO V., POHL T.M., PORTEFELLE D., PORWOLIK S., PRESCOTT A.M.,  
 RA PRESECAN E., PUJIC P., PURNELLE B., RAPORT G., REY M., REYNOLDS S.,  
 RA RIEGER M., RIVOLTA C., ROCHA E., ROCHE B., ROSE M., SADIE Y.,  
 RA SATO T., SCALIANI E., SCHLEITCH S., SCHROETER R., SCOFFONE F.,  
 RA SEKIUCHI J., SERKOWSKA A., SEROR S.J., SERROR P., SHIN B.S., SOLDO B.,  
 RA SOROKIN A., TACCONI E., TAKAGI T., TAKAHASHI H., TAKEMARU K.,  
 RA TAKEUCHI M., TAMAKOSHI A., TANAKA T., TERNSTRA P., TOGNONI A.,  
 RA TOSATO V., UCHIYAMA S., VANDENBOL M., VANNIER F., VASSAROTTI A.,  
 RA VIARI A., WAMBUTT R., WEDLER E., WETZENBGER T.,  
 RA WINTERS P., WIPAT A., YAMAMOTO H., YAMANE K., YASUMOTO K.,  
 RA YOSHIDA K., YOSHIKAWA H.F., ZUMSTEIN E., YOSHIKAWA H., DANCHIN A.,  
 RT "The complete genome sequence of the gram-positive bacterium *Bacillus subtilis*";  
 RT  
 RL Nature; 390:249-256(1997).  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=168;  
 RA KUNST F., OGASAWARA N., YOSHIKAWA H., DANCHIN A.;  
 RC Submitted (Nov-1997) to the EMBL/GenBank/DBJ databases.  
 DR EMBL; Z99110; CAB12998; 1; -.  
 SQ SEQUENCE 250 AA; 30.119 MW; C96222FD CRC32;

Query Match 83.9%; Score 47; DB 2; Length 250;  
 Best Local Similarity 66.7%; Pred. No. 1.0e+01;  
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Db 89 TDGIGILAV 97  
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 QY 1 AGIGILTV 9

RESULT 5 PRELIMINARY; PRT; 478 AA.  
 ID 037090; AC 037090; DT 01-NOV-1996 (TREMBLrel. 01, Created)  
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
 DE GLYCOPROTEIN GII.  
 OS Pseudorabies virus.  
 OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;  
 OC Alphaherpesvirinae; Varicellovirus.  
 RN 111  
 RP SEQUENCE FROM N.A.  
 RC SPAIN=INDIANA S;  
 RX MEDLINE; 96316347.  
 RA ISHIKAWA K., TSUTSUI M., TAGUCHI K., SAITO A., MURAMATSU M.;

RT "Sequence variation of the gc gene among pseudorabies virus strains.";  
 RL Vet. Microbiol. 49:267-272(1996).  
 DR EMBL: D49436; BAA08A14.1;  
 PRINTS: PR00668; GLYCOPROTEIN.C.  
 SQ SEQUENCE 478 AA; 51150 MW; D6A143B4 CRC32;

Query Match Score 47; DB 14; Length 478;  
 Best Local Similarity 75.0%; Pred. No. 1.01e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Db 455 AGIGILAI 462  
 |||||:  
 Qy 2 AGIGILTV 9

RESULT 6 PRELIMINARY; PRT; 479 AA.  
 ID Q87089; PRELIMINARY; PRT; 479 AA.  
 AC 087089;  
 DT 01-NOV-1996 (TREMBLrel. 01, Created)  
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)  
 DE GLYCOPROTEIN GI11.  
 OS Pseudorabies virus.  
 OC dsDNA viruses, no RNA stage; Herpesviridae;  
 OC Alphaherpesvirinae; Varicellovirus.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-YAMAGATA S-81;  
 RX 96316347.  
 RA ISHIKAWA K., TSUTSUI M., TAGUCHI K., SAITO A., MURAMATSU M.,  
 RT "Sequence variation of the gc gene among pseudorabies virus strains.";  
 RL Vet. Microbiol. 49:267-272(1996).  
 DR EMBL: D49435; BAA08A13.1;  
 PRINTS: PR00668; GLYCOPROTEIN.C.  
 SQ SEQUENCE 479 AA; 51109 MW; A009EB9B CRC32;

Query Match Score 47; DB 14; Length 479;  
 Best Local Similarity 75.0%; Pred. No. 1.01e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Db 456 AGIGILAI 463  
 |||||:  
 Qy 2 AGIGILTV 9

RESULT 7 PRELIMINARY; PRT; 479 AA.  
 ID Q87091; PRELIMINARY; PRT; 479 AA.  
 AC 087091;  
 DT 01-NOV-1996 (TREMBLrel. 01, Created)  
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)  
 DE GLYCOPROTEIN GI11.  
 OS Pseudorabies virus.  
 OC dsDNA viruses, no RNA stage; Herpesviridae;  
 OC Alphaherpesvirinae; Varicellovirus.  
 RN [1]  
 RP SEQUENCE FROM N.A.

RC MEDLINE: 96316347.  
 RA ISHIKAWA K., TSUTSUI M., TAGUCHI K., SAITO A., MURAMATSU M.,  
 RT "Sequence variation of the gc gene among pseudorabies virus strains.";  
 RL Vet. Microbiol. 49:267-272(1996).  
 DR EMBL: D49437; BAA08A15.1;  
 PRINTS: PR00668; GLYCOPROTEIN.C.  
 SQ SEQUENCE 479 AA; 51148 MW; CC3EEFF9A CRC32;

Query Match Score 47; DB 14; Length 479;  
 Best Local Similarity 75.0%; Pred. No. 1.01e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Db 456 AGIGILAI 463  
 |||||:  
 Qy 2 AGIGILTV 9

RESULT 8 PRELIMINARY; PRT; 848 AA.  
 ID O18139; PRELIMINARY; PRT; 848 AA.  
 AC O18139;  
 DT 01-JAN-1998 (TREMBLrel. 05, Created)  
 DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)  
 DT 01-JAN-1999 (TREMBLrel. 09, Last annotation update)  
 DE T26H2.7 PROTEIN.  
 GN T26H2.7  
 OS Caenorhabditis elegans.  
 OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditida;  
 OC Rhabditina; Rhabditoidae; Rhabditidae; Pelederinae; Caenorhabditida.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RA MATTHEWS L.;  
 RA Submitted (NOV-1996) to the EMBL/GenBank/DDBJ databases.  
 RN [2]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE; 94150718.  
 RA WILSON R., ALNSCOUGH R., ANDERSON K., BAYNES C., BERKS M., COOPER J., COULSON A., FULTON L., GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L., JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P., LIGHTNING J., LLOYD C., MC MORRAY B., MORTMORE B., O'CALLAGHAN M., PARSONS J., PERCY C., RIEKEN L., ROOPRA A., SAUNDERS D., SHONNIEEN R., SHALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J., THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R., WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;  
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C. elegans.";  
 RL Nature 368:32-38(1994).  
 DR EMBL: 282055; CAB04848.1;  
 SEQ SEQUENCE 848 AA; 98312 MW; 371853A7 CRC32;

Query Match Score 46; DB 5; Length 848;  
 Best Local Similarity 66.7%; Pred. No. 1.67e+01;  
 Matches 6; Conservative 3; Mismatches 0; Indels 0; Gaps 0;  
 Db 340 SASIGILTV 348  
 :|||||:  
 Qy 1 AAGIGILTV 9

RESULT 9 PRELIMINARY; PRT; 339 AA.  
 ID O30640; PRELIMINARY; PRT; 339 AA.  
 AC O30640;  
 DT 01-JAN-1998 (TREMBLrel. 05, Created)  
 DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)  
 DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)  
 DE METHYLCOBAMIDE:COM METHYLTRANSFERASE ISOZYME A.  
 GN MBTA.  
 OS Methanosaרכינה barkeri.  
 OC Archaea; Euryarchaeota; Methanosaרכינae; OC Methanosaרכינה.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN-M (DSM 800);  
 RX MEDLINE; 97341199.  
 RA BURKE S.A., KRZYCZKI J.A.;  
 RT "Reconstitution of Monomethylamine:Coenzyme M methyl transfer with a corrinoid protein and two methyltransferases purified from a Methanosaרכינה barkeri.";  
 RT JL Biol. Chem. 272:16370-16577(1997).  
 DR EMBL: AF013713; AAC38632.1;  
 DR PDB: P01208; URO-D;  
 KW Transferase; Methyltransferase.  
 SQ SEQUENCE 339 AA; 36664 MW; 040E3CF3 CRC32;

Query Match Score 45; DB 1; Length 339;  
 Best Local Similarity 75.0%; Pred. No. 2.73e+01;  
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Db 456 AGIGILAI 463  
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 Qy 2 AGIGILTV 9

Db 307 AGYGLTV 314  
Qy 2 AGIGILTV 9

RESULT 12  
ID 048172; PRELIMINARY; PRT; 370 AA.  
AC 048172;  
AC 048172; 01-JUN-1998 (TREMBLrel. 06, Created)  
DT 01-JUN-1998 (TREMBLrel. 06, Last sequence update)  
DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)  
DE CYTOCHROME B.  
GN CCB.  
OS Polytomella sp. 'Pringsheim' 198.80%/  
OS Mitochondrion.  
OC Viridiplanteae; Chlorophyta; Chlorophyceae; Volvocales;  
OC Chlamydomonadaceae; Polytomekia  
RN [1]  
RN RP SEQUENCE FROM N.A.  
RC STRAIN=198.80; FROM E.G.; PRINGSHEIM;  
RA ANTARAMIAN A.; FUNES ARGUELLO S.; VASQUEZ ACEVEDO M.; CORIA R.;  
RA GONZALEZ-HALPHEN D.;  
RL Submitted (JAN-1997) to the EMBL/GenBank/DDBJ databases.  
DR U87396; AAC24896.1; -  
DR MENDEL; 23385; Pols; cob; 23385.  
DR PFAM; PF00022; cytochrome\_b\_C; 1;  
DR PEAM; PF00033; cytochrome\_b\_N; 1;  
KW Mitochondrion.  
SQ SEQUENCE 370 AA; 5D617081 CRC32;

Query Match 80.4%; Score 45; DB 8; Length 370;  
Best Local Similarity 85.7%; Pred. No. 2.73e+01;  
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;  
Db 288 GIGILAV 294  
Qy 3 GIGILTV 9

RESULT 13  
ID Q63276; PRELIMINARY; PRT; 420 AA.  
AC Q63276;  
AC 01-NOV-1996 (TREMBLrel. 01, Created)  
AC 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
DT 01-JAN-1999 (TREMBLrel. 09, Last annotation update)  
DE KAN-1.  
OS Rattus norvegicus (Rat).  
OC Eutheria; Rodentia; Sciurognathida; Muridae; Murinae; Rattus.  
RN [1]  
RP SEQUENCE FROM N.A.  
RC STRAIN=TWISTER; TISSUE=LIVER;  
RX MEDLINE; 96003917.  
RA FURUTANI M.; ARII S.; HIGASHITSUJI H.; MISE M.; FUKUMOTO M.;  
RA TAKANO S.; NAKAYAMA H.; IIMURA M.; FUJITA J.;  
RT "Reduced expression of kan-1 (encoding putative bile acid-CoA-amino  
acid N-acyltransferase) mRNA in livers of rats after partial  
hepatectomy and during sepsis.";  
RL Biochem J; 311:203-208(1995).  
DR BAA07901.1;  
SQ SEQUENCE 420 AA; 46496 MW; 7B62AACF CRC32;

Query Match 80.4%; Score 45; DB 11; Length 420;  
Best Local Similarity 55.6%; Pred. No. 2.73e+01;  
Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;  
Db 226 GPGVGILSV 234  
Qy 1 AAGIGILTV 9

RESULT 14  
ID 008833; PRELIMINARY; PRT; 420 AA.  
AC 008833;  
DT 01-JUL-1997 (TREMBLrel. 04, Created)  
DT 01-JUL-1997 (TREMBLrel. 04, Last sequence update)

Db 307 AGYGLTV 314  
Qy 2 AGIGILTV 9

Query Match 80.4%; Score 45; DB 1; Length 339;  
Best Local Similarity 75.0%; Pred. No. 2.73e+01;  
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)  
DE BILE ACID COA: AMINO ACID N-ACYLTRANSFERASE.  
GN BAAT.  
OS Mus musculus (Mouse).  
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;  
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
RN [1]  
RP SEQUENCE FROM N.A.  
RA FALANY C.N.; FORTINBERRY H.; LEITER E.H.; BARNES S.;  
RL J. Lipid Res. 0:0(0);  
DR U95215; AAB5825.1; -.  
DR MGI:106642; Biot.  
KW Transferase; Acyltransferase.  
SEQUENCE 420 AA; 46528 MW; 4A22EFFC CRC32;  
SQ :||:|||||:  
Db 225 GPNGLSLV 233  
QY 1 AAGIGILTV 9

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RESULT 15  
ID Q17592 PRELIMINARY; PRT; 980 AA.  
AC Q17592;  
DT 01-NOV-1996 (TREMBLrel. 01, Created)  
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)  
DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)  
DE SIMILARITY TO INSULIN-DEGRADING ENZYMES.  
GN C0265..1.  
OS Caenorhabditis elegans.  
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditida;  
OC Rhabdita; Rhabditoidea; Peledorinae; Caenorhabditis.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX STRAIN=BRISTOL N2;  
RA MEDLINE: 94150718  
RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M.,  
RA COOPER J., COULSON A., COUPPEY T., CORNELL M., COPSEY T., COOPER J., COULSON A.,  
RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,  
RA GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L.,  
RA JONES M., KERSHAW J., KIRSTEN N., LAISTER N., LATTEILLE P.,  
RA LIGHTNING J., LLOYD C., MCCLURAY A., MORTIMORE B., O'CALLAGHAN M.,  
RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,  
RA SMALDON N., SMITH A., SONNHAMMER E., SPADON R., SULSTON J.,  
RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,  
RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLFARTH P.;  
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.  
RT elegans";  
RL Nature 368:32-38 (1994).  
RN [2]  
RP SEQUENCE FROM N.A.  
RC STRAIN=BRISTOL N2;  
RA BENTLEY D., KEMP K., SCHEET P.; Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.  
RN [3]  
RP SEQUENCE FROM N.A.  
RC STRAIN=BRISTOL N2;  
RA WATERSTON R.; Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.  
RL U55372; AA949001.1;  
DR PROTE; PS00143; INSULINASE; 1.  
DR PF00675; Peptidase\_M16; 1.  
SQ SEQUENCE 980 AA; 6D56C08D CRC32;  
Db 303 AAGFGILNV 311  
||:|||||:  
QY 1 AAGIGILTV 9  
Search completed: Fri May 5 22:03:19 2000  
Job time : 89 secs.

(TM)

Release 3.1A John F. Collins, Biocomputing Research Unit.  
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 protein - protein database search, using Smith-Waterman algorithm  
 on on: Fri May 5 22:00:53 2000; MasPar time 3.16 Seconds  
 tabular output not generated.  
 85.040 Million cell updates/sec

Title: >US-09-267-439-4  
Description: (1-9) from US09267439.pep  
Aspect, Score: 56

scoring table:  
PAM 150  
Gap 15

searched: 82229 seqs, 29864866 residues  
estimated processing: Minimum Match %

post processing. Listing first 45 summaries

database: swissprot  
swissprot:1

Statistics: Mean 22.670; Variance 23.392; scale 0.969

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	ID	Description	Pred. No.
------------	-------	-------------	--------	----	-------------	-----------

1 56 100.0 118 1 MARI HUMAN MELANOMA RECOG 1.80e-02  
 2 47 83.0 101 1 MARI SITI AC MEMBRANE ASSOCIATED 2.15e-02  
 3

2	*,*	82.2	1	MELANOMA ASSOCIATED AI	3.15e+00
3	47	83.9	479	1 VOGLC PROVIF GLYCOPROTEIN GIII PREC	
4	46	82.1	231	1 YSA1 YEAST YSA1 PROTEIN.	5.37e+00

5	45	80.4	291	1	YATQ RHIISN PROBABLE PEPTIDE ABC T	9.09e+00
6	44	78.6	394	1	FTS2_AAOV1	1.52e+01
7	44	78.6	456	1	GIMT1_ECOLI	1.52e+01
					HDP-N-ACETYLGLUCOSAMIN	

8	44	78.6	635	1	XYND_PAEPO	ENDO-1,4-BETA-XYLANASE	1.52e+01
9	43	76.8	132	1	ATPE_ARATH	ATP SYNTHASE EPSILON C	2.53e+01
10	43	76.8	132	1	ATPE_ARATH	ATP SYNTHASE EPSILON C	2.53e+01
11	43	76.8	132	1	ATPE_ARATH	ATP SYNTHASE EPSILON C	2.53e+01
12	43	76.8	132	1	ATPE_ARATH	ATP SYNTHASE EPSILON C	2.53e+01

10	4.3	/6.8	444	1
11	4.3	76.8	461	1
12	4.3	76.8	493	1

14	4.3	76.8	611	1	YD3M_HERAU	HYPOTHETICAL
14	4.2	75.0	110	1	YD3M_MBTVO	HEAD DECORATION PROTEI
15	4.2	75.0	216	1	YF1A2_MFTVO	FLAGELLIN B2 DEPPENTSCOP
						A_15E+01

16	4.2	75.0	218	1	FLAGELLIN B1 PRECURSOR
17	4.2	75.0	222	1	FLAGELLIN B2 PRECURSOR

18	4.2	75.0	308	1	MENA_HAEIN	1,4-DIHYDROXY-2-NAPHTH	4.15e+01
19	4.2	75.0	325	1	RCEM_CHRV1	REACTION CENTER PROTEIN	4.15e+01
20	4.2	75.0	3332	1	ACOA_ALC1	ACETOIN 2,6-DICHOROP	4.15e+01

21	42	75.0	345	1
22	42	75.0	461	1
23	42	75.0	601	1
				POSSIBLE THIOPHENE AND POSSIBLY THIOPHENYL SUBSTITUTED BENZENE

24	4.2	75..0	503	1	SECD...HELPY	PROTEIN EXPORT MEMBRAN	4.15e+01
25	4.2	75..0	526	1	HELPY	PROTEIN EXPORT MEMBRAN	4.15e+01
26	4.2	75..0	530	1	AIP2...YEAST	ACTIN INTERACTING PROT	4.15e+01
27	4.2	75..0	659	1	YYBT...BACSU	HYPOTHETICAL 74.3 KD P	4.15e+01
28	4.2	75..0	885	1	YDGH...BACSU	PUTATIVE MEMBRANE PROT	4.15e+01
29	4.2	75..0	1325	1	YDEK...ECOLI	HYPOTHETICAL 136.5 KD	4.15e+01
30	4.2	75..0	1530	1	BFR1...SCHPO	BREFELDIN A RESISTANCE	4.15e+01
31	4.1	73..2	225	1	CD9	FEELCA	6.75e+01
32	4.1	73..2	237	1	BACT...HALSA	SENSORY RHODOPSIN II {	6.75e+01
33	4.1	73..2	271	1	KY23...	HYPOTHETICAL 31.0 KD P	6.75e+01
34	4.1	73..2	313	1	YATP...RHISN	PROBABLE PEPTIDE ABC T	6.75e+01
35	4.1	73..2	337	1	OPSX...HUMAN	VISUAL PIGMENT-LIKE RE	6.75e+01
36	4.1	73..2	359	1	HYD...CORG1	HYPOTHETICAL PROTEIN I	6.75e+01
37	4.1	73..2	384	1	PQOE...METEX	COENZYME PQO SYNTHESIS	6.75e+01
38	4.1	73..2	401	1	YABA...SCHPO	HYPOTHETICAL 44.4 KD P	6.75e+01
39	4.1	73..2	487	1	Y346...	HYPOTHETICAL 52.2 KD P	6.75e+01
40	4.1	73..2	633	1	Y561...HAEIN	HYPOTHETICAL PROTEIN H	6.75e+01
41	4.1	73..2	666	1	Y561...BRAJA	PROBABLE CYTOCHROME C	6.75e+01
42	4.1	73..2	845	1	MAT3...RAT	MATRIN 3.	6.75e+01
43	4.1	73..2	977	1	YDG8...SCHPO	HYPOTHETICAL 111.4 KD	6.75e+01
44	4.1	73..2	1109	1	CYAB...CANFA	RETINAL GUANYL CYCLAT	6.75e+01
45	4.1	73..2	1331	1	CYAB...LEILO	RECEPTOR-TYPE ADENYLAT	6.75e+01

Query Match	100.0%	Score 56;	DB 1;	Length 118;		
Best Local Similarity	100.0%	Pred. No.	1.80e-02;			
Matches	9;	Conservative	0;	Indels	0;	Gaps 0;
DR	27 AGIGILTV 35					
QY	• 1 AGIGILTV 9					
RESULT	2	ATPL_SULAC	STANDARD;	PRT;	101 AA.	
ID	ATPL_SULAC					
AC	P2340;					
DT	01-NOV-1991 (Rel. 20, Created)					
DT	01-NOV-1991 (Rel. 20, Last sequence update)					
DT	15-JUL-1998 (Rel. 36, Last annotation update)					
DE	MEMBRANE-ASSOCIATED ATPASE C CHAIN (EC 3.6.1.34) (SUL-ATPASE DE PROTEOLIPID CHAIN).					
GN	ATPP.					
OS	Sulfolobus acidocaldarius.					
OC	Archaea; Crenarchaeota; Sulfolobales; Sulfolobus.					
RN	[1]					
RP	SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.					
RX	MEDLINE: 89214142.					
RA	DENDA K., KONISHI J., OSHIMA T., DATE T., YOSHIDA M.;					
RT	"A gene encoding the proteolipid subunit of <i>Sulfolobus acidocaldarius</i> ATPase complex";					
RT	ATPase complex.					
RL	J. Biol. Chem. 264: 7119-7121(1989).					
CC	-1- FUNCTION: THE C CHAIN IS A PROTEOLIPID, AND ONE OF THE MEMBRANOUS SUBUNITS OF THE NONENZYMATIC COMPONENT OF THE SUL-ATPASE COMPLEX.					
CC	-1- SUBUNIT: SUL-ATPASE IS COMPOSED OF SIX (OR FIVE?) SUBUNITS:					
CC	ALPHA, BETA, DELTA, GAMMA, C (PROTEOLIPID), AND POSSIBLY EPSILON.					
CC	-1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POENTIAL).					
CC	-1- SIMILARITY: BELONGS TO THE V-ATPASE PROTEOLIPID SUBUNIT FAMILY.					
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CC	DR	J04740; AAA72703.1;	-			
DR	PIR; A33351; A33351.					
DR	HSSP; P00138; ICGN.					
DR	PFAM; PF00137; ATP-synt_C; 1.					
KW	Hydrogen ion transport; Lipid-binding; Transmembrane.					
FT	TRANSMEM 5 25 POTENTIAL.					
FT	TRANSMEM 37 57 POTENTIAL.					
FT	TRANSMEM 75 95 POTENTIAL.					
SQ	SEQUENCE 101 AA: 10362 MW: 1DC8C74D CRC32;					
Query Match	83.9%	Score 47;	DB 1;	Length 101;		
Best Local Similarity	87.5%	Pred. No.	3.15e+00;			
Matches	7;	Conservative	0;	Indels	0;	Gaps 0;
DR	59 AGIGILTV 66					
QY	• 1 AGIGILT 8					
RESULT	3	VGLC_PRVIF	STANDARD;	PRT;	479 AA.	
ID	P06724;					
DT	13-AUG-1987 (Rel. 05, Created)					
DT	13-AUG-1987 (Rel. 05, Last sequence update)					
DT	01-APR-1993 (Rel. 25, Last annotation update)					
DE	GLYCOPROTEIN GTII PRECURSOR.					
OS	Pseudorabies virus (strain Indiana-Funkhauser / Becker) (PRV).					
OC	Viruses; dsDNA viruses, no RNA state; Herpesviridae;					
OC	Alphaherpesvirinae; Varicellovirus.					
RN	[1]					
RP	SEQUENCE FROM N.A.					
RX	MEDLINE: 86200375.					
RA	ROBBINS A.K., WATSON R.J., WHEALY M.E., HAYS W.W., ENQUIST L.W.;					
RT	"Characterization of a pseudorabies virus glycoprotein gene with homology to herpes simplex virus type 1 and type 2 glycoprotein C.";					
RL	J. Virol. 55:339-347(1986).					
CC	-1- SIMILARITY: TO OTHER HERPESVIRUSES GLYCOPROTEIN C.					
CC	-1- SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN GENE SUPERFAMILY.					
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CC	DR	M12778; AAA7464.1;	-			
DR	PIR; A26097; VGBEPB.					
KW	Glycoprotein; Transmembrane; Signal.					
FT	SIGNAL 1 22 GLYCOPROTEIN GTII.					
FT	CHAIN 23 479 GLYCOPROTEIN GTII.					
FT	CARBOHYD 40 40 POTENTIAL.					
FT	CARBOHYD 84 84 POTENTIAL.					
FT	CARBOHYD 169 169 POTENTIAL.					
FT	CARBOHYD 192 192 POTENTIAL.					
FT	CARBOHYD 220 220 POTENTIAL.					
FT	CARBOHYD 228 228 POTENTIAL.					
FT	CARBOHYD 285 285 POTENTIAL.					
FT	CARBOHYD 302 302 POTENTIAL.					
SQ	SEQUENCE 479 AA: 51206 MW: 42EE5703 CRC32;					
Query Match	83.9%	Score 47;	DB 1;	Length 479;		
Best Local Similarity	75.0%	Pred. No.	3.15e+00;			
Matches	6;	Conservative	0;	Mismatches 2;	Indels 0;	Gaps 0;
Db	456 AGIGILAI 463					
Db	: 2 AGIGILTV 9					
QY						
RESULT	4	YSA1_YEAST	STANDARD;	PRT;	231 AA.	
ID	YSA1_YEAST					
AC	Q01976;					
DT	01-OCT-1993 (Rel. 27, Created)					
DT	01-OCT-1994 (Rel. 30, Last sequence update)					
DE	YSA1 PROTEIN.					
GN	YSA1 OR YBR11C OR YBR0907.					
OS	Saccharomyces cerevisiae (Baker's yeast).					
OC	Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales.					
RN	[1]					
RP	SEQUENCE FROM N.A.					
RC	STRAIN=S288C;					
RC	MEDLINE: 95208357.					
RX	"Analysis of a 70 kb region on the right arm of yeast chromosome II."					
RA	MANNHAUPT G., STUCKA R., EHNL S., VETTER I., FELDMANN H.; Yeast 10:1363-1381(1994).					
RN	[2]					
RP	SEQUENCE OF 1-47 FROM N.A.					
RC	STRAIN=S288C;					
RX	MEDLINE: 92327848.					
RA	MANNHAUPT G., STUCKA R., EHNL S., VETTER I., FELDMANN H.;					
RT	"Molecular analysis of yeast chromosome II between CMD1 and LYS2: the excision repair gene Rad16 located in this region belongs to a novel group of double-finger proteins."					
RL	Yeast 8:397-408(1992).					
CC	-1- SIMILARITY: STRONG, TO B. SUBTILIS YQKG.					
CC	-1- SIMILARITY: TO PROTEINS WITH A CORE MUTT DOMAIN.					
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CC	or send an email to <a href="mailto:license@isb-sib.ch">license@isb-sib.ch</a> ).
CC	-----
DR	Z35580; CAA85068.1;
DR	X78993; CAA55614.1;
DR	X66247; CAA46972.1;
PIR	S44631; S44691.
DR	L0002551; YSA1.
DR	PROSITE: PS00893; M0UTT; 1.
DR	PFAM: PF00293; mutR; 1.
DOMAIN	112 145 MOTU-LIKE.
SO	231 AA; 26087 MW; 49A2D6CB CRC32;
Query Match	82.1%
Best Local Similarity	85.7%
Matches	6;
Conservative	1;
Mismatches	0;
Indels	0;
Gaps	0;
Score	46;
DB	DB 1;
Length	231;
Pred. No.	5.37e+00;
FT	
SEQUENCE	231 AA;
Db	79 GIGILTI 85
Qy	: 85
3 GIGILTV 9	
RESULT	5
ID	Y4TQ_RHISN
STANDARD	
PRT	291 AA.
AC	Q53192;
DT	01-NOV-1997 (Rel. 35, Created)
DT	01-NOV-1997 (Rel. 35, Last sequence update)
DT	01-NOV-1997 (Rel. 35, Last annotation update)
DE	PROTEIN PEPTIDE ABC TRANSPORTER PERMEASE PROTEIN Y4TQ.
GN	Y4TQ.
OS	Rhizobium sp. (strain NGR34).
OG	Plasmid symb phGR34a.
OC	Bacteriia; Proteobacteria; alpha subdivision; Rhizobiaceae group;
OC	Rhizobiaceae; Rhizobium.
RN	[1]
RP	SEQUENCE FROM N.A.
RX	MEDLINE; 9730556.
RX	FREIBERG C., PERRET X., BROUGHTON W.J., ROSENTHAL A.,
RA	FREEBERG C.A., FELLAY R., BAIROCH A., BROUGHTON W.J., ROSENTHAL A.,
RA	PERRET X.,
RT	"Molecular basis of symbiosis between Rhizobium and legumes.";
RL	Nature 387:394-401(1997).
[2]	
RP	SEQUENCE FROM N.A.
RX	MEDLINE; 96389014.
RX	FREIBERG C., PERRET X., BROUGHTON W.J., ROSENTHAL A.,
RT	"Sequencing the 500-kb GC-rich symbiotic replicon of Rhizobium sp. NGR34 using dye terminators and a thermostable 'sequenase': a beginning."
RT	beginning."
RL	Genome Res. 6:590-600(1996).
-I	FUNCTION: PROBABLY PART OF A BINDING-PROTEIN-DEPENDENT TRANSPORT SYSTEM Y4TOPS FOR A PEPTIDE. PROBABLY RESPONSIBLE FOR THE TRANSLLOCATION OF THE SUBSTRATE ACROSS THE MEMBRANE.
-I	SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. INNER MEMBRANE (POTENTIAL).
-I	SIMILARITY: WITH INTEGRAL MEMBRANE COMPONENTS OF OTHER BINDING-PROTEIN-DEPENDENT TRANSPORT SYSTEMS. BELONGS TO THE OPPBC SUBFAMILY.
CC	-----
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CC	or send an email to <a href="mailto:license@isb-sib.ch">license@isb-sib.ch</a> ).
CC	-----
DR	Z68203; CAA92399.1;
DR	AAB91870.1;
DR	PROSITE: PS00098; BPD_TRANS_INN_MEMBR; 1;
DR	PFAM: PF00528; BPD_transp_1.

KW	Hypothetical protein; Transport; Amino-acid transport; Transmembrane;
KW	Inner membrane; Plasmid.
FT	TRANSEM 28 48 POTENTIAL.
FT	TRANSEM 92 112 POTENTIAL.
FT	TRANSEM 137 213 233 POTENTIAL.
FT	TRANSEM 249 269 POTENTIAL.
FT	TRANSEM 291 AA; 30910 MW; 3263271E CRC32;
SEQUENCE	
Query Match	Score 45; DB 1; Length 291;
Best Local Similarity	66.7%; Pred. No. 9.09e+00;
Matches	6; Conservative 2; Mismatches 1; Indels 0; Gaps
Db	147 GPGIGILIV 155
Qy	:        1 AAGIGILTV 9
RESULT	6
ID	FTSZ_AZOVI STANDARD; PRT; 394 AA.
AC	P77817;
DT	01-NOV-1997 (Rel. 35, Created)
DT	01-NOV-1997 (Rel. 35, Last sequence update)
DT	15-DEC-1999 (Rel. 39, Last annotation update)
DE	CELL DIVISION PROTEIN FT SZ.
GN	FTSZ.
OS	Azotobacter vinelandii
OC	Bacteria; Proteobacteria; gamma subdivision; Azotobacteraceae;
OC	Azotobacter.
RN	[1]
RP	SEQUENCE FROM N.A., AND CHARACTERIZATION.
RC	STRAIN=DJ116;
RX	MEDLINE; 982267010.
RA	LU C., STRICKER J., ERICKSON H.P.;
RT	"Ftsz from Escherichia coli, Azotobacter vinelandii, and Thermotoga maritima -- quantitation, GTP hydrolysis, and assembly.";
RL	Cell Motil. Cytoskeleton 40:71-86 (1998).
CC	-!- FUNCTION: THIS PROTEIN IS ESSENTIAL TO THE CELL-DIVISION PROCESS. IT'S SEEMS TO ASSEMBLE INTO A DYNAMIC RING ON THE INNER SURFACE OF THE CITOPLASMIC MEMBRANE AT THE PLACE WHERE DIVISION WILL OCCUR, AND THE FORMATION OF THE RING IS THE SIGNAL FOR SEPTATION TO BEGIN. BINDS TO AND HYDROLYSES GTP.
CC	-!- SUBUNIT: AGGREGATE TO FORM A RING-LIKE STRUCTURE.
CC	-!- SUBCELLULAR LOCATION: CITOPLASMIC. ASSEMBLE AT THE INNER SURFACE OF THE CITOPLASMIC MEMBRANE (BY SIMILARITY).
CC	-!- SIMILARITY: BELONGS TO THE FTSZ FAMILY.
CC	
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DR	EMBL; U65339; AAC24603.1; -.
DR	HSSP; Q57816; IFSZ.
DR	PROSITE; PS01134; FTSZ_1; 1.
DR	PROSITE; PS01135; FTSZ_2; 1.
DR	Cell division; Septation; GTP-binding.
KW	NP_BIND 104 112 GTP (POTENTIAL).
FT	SEQUENCE 394 AA; 41153 MW; 4EB8134 CRC32;
SQ	
Query Match	Score 44; DB 1; Length 394;
Best Local Similarity	77.8%; Pred. No. 1.52e+01;
Matches	7; Conservative 1; Mismatches 1; Indels 0; Gaps
Db	121 AKGLGILTV 129
Qy	:      1 AAGIGILTV 9
RESULT	7

ID GLMU\_ECOLI STANDARD; PRT; 456 AA.  
AC P17114; P76746;  
DT 01-AUG-1990 (Rel. 15, Created)  
DT 01-NOV-1997 (Rel. 35, Last sequence update)  
DT 01-NOV-1997 (Rel. 35, Last annotation update)  
DE UDP-N-ACETYLGLUCOSAMINE PYROPHOSPHORYLASE (EC 2.7.7.23) (N-  
DE GLMU.  
GN Escherichia coli.  
OS Escherichia coli.  
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;  
OC Escherichia.  
RN [1]  
RP SEQUENCE FROM N.A.  
RX MEDLINE: 85121806;  
RA WALKER J.E., GAY N.J., SARASTE M., EBERLE A.N.;  
RT "DNA sequence around the Escherichia coli unc operon. Completion of  
RT the sequence of a 17 kilobases segment containing asnA, oricC, unc,  
RT gms and phoS.";  
RT Biochem. J. 224:799-815(1984).  
RN [2]  
RP SEQUENCE FROM N.A.  
RX STRAIN-K12 / MG1655;  
RA BURLAND V.D., PLUNKETT G. III, DANIELS D.L., BLATTNER F.R.;  
RT "DNA sequence and analysis of 136 kilobases of the Escherichia coli  
RT genome: organizational symmetry around the origin of replication."  
RN [3]  
RP REVISIONS.  
RX STRAIN-K12 / MG1655;  
RA BLATTNER F.R., PLUNKETT G. III, BLOCH C.A., PERNA N.T., BURLAND V.,  
RA RILEY M., COLLADO-VIDES J., GLASNER J.D., RODE C.K., MAYHEW G.F.,  
RA RILEY M., COLLADO-VIDES J., GLASNER F.D., RODE C.K., MAYHEW G.F.,  
RA MAU B., SHAO Y., KIRKPATRICK H.A., GOEDEN M.A., ROSE D.J.,  
RA "The complete genome sequence of Escherichia coli K-12.";  
RT Science 277:1453-1474 (1997).  
RN [4]  
RP IDENTIFICATION.  
RX MEDLINE: 94012475.  
RA MENGIN-LEGREUL D., VAN HEIJENOORT J.;  
RT "Identification of the glmU gene encoding N-acetylglucosamine-1-  
RT phosphate uridyltransferase in Escherichia coli.";  
RL J. Bacteriol. 175:6150-6157 (1993).  
CC -1- FUNCTION: BIFUNCTIONAL ENZYME RESPONSIBLE FOR THE ACETYLATION OF  
CC GLC-N-1-P TO GIVE GLCNAC-1-P AND THE SYNTHESIS OF UDP-GLCNAC.  
CC -1- CATALYTIC ACTIVITY: UDP + N-ACETYL-ALPHA-D-GLUCOSAMINE  
CC -1- PHOSPHATE = PYROPHOSPHATE + UDP-N-ACETYL-ALPHA-D-GLUCOSAMINE.  
CC -1- PATHWAY: PEPTIDOGLYCAN AND LIPOPOLYSACCHARIDE BIOSYNTHESIS.  
CC -1- SIMILARITY: BELONGS TO THE CYSTEYLACYLIPXA/ANODL FAMILY OF  
CC ACETYLTRANSFERASES. COMPOSED OF MULTIPLE REPEATS OF [LIV]-G-X-(4).  
CC -1- CAUTION: REF. 2 SEQUENCE DIFFERS FROM THAT SHOWN DUE TO A  
CC FRAMESHIFT THAT CREATES TWO ORFS.  
CC  
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CC  
DR X57094; CAA040378.1;  
CC Xylan degradation; Hydrolase; Glycosidase; Signal.  
FT FT SIGNAL  
FT CHAIN  
SQ SEQUENCE 635 AA; 67914 MW; 078AAB82 CRC32;

Query Match Score 44; DB 1; Length 635;  
Best Local Similarity 75.0%; Pred. No. 1.52e+01;  
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;  
Db 149 GAGIGVLT 156  
Qy 1 AAGIGILT 8

RESULT 9  
ID ATP\_E\_ARATH STANDARD; PRT; 132 AA.  
AC PD9468;  
DT 01-MAR-1989 (Rel. 10, Created)  
DT 01-MAR-1999 (Rel. 10, Last sequence update)  
DT 01-OCT-1994 (Rel. 30, Last annotation update)  
DE ATP SYNTHASE EPSILON CHAIN (EC 3.6.1.34).  
GN ATP.  
OS Arthropods.  
OG Chloroplast.  
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; tracheophyta;

EMBL; X01631; CAA25784.1;  
DR EMBL; L10328; AAA62082.1; ALT\_FRAME.  
DR EMBL; L10328; AAA62081.1; ALT\_FRAME.  
DR AE000450; AAC176753.1; -.  
ECOGENE: EG11198; GLMU.  
DR PROST1; PS00101; HEXAPEP\_TRANSFERASE; 1.  
DR PFAM; PF00132; hexapep\_transferase; 3.  
DR PFAM; PF00483; NTP\_transferase; 1.  
DR Peptidoglycan synthesis; Cell wall; transferase;  
KW Nucleotidyltransferase; Repeat; Multifunctional enzyme.  
FT CONFLICT 186 187 KL -> NV (IN REF. 1).

OC euphyllophytes; Spermatophyta; Magnoliophyta; eudicotsyledons;  
 OC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;  
 OC Arabidopsis.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=CV: LANDSBERG ERECTA;  
 RX MEDLINE: 89057486.  
 RA CHEN H.-C., WINTZ H., WELL J.-H., PILLAY D.T.N.;  
 RT "Nucleotide sequence of chloroplast CF1-ATPase epsilon-subunit and  
 elongator tRNA<sub>Met</sub> genes from Arabidopsis thaliana.";  
 RL Nucleic Acids Res. 16:1032-1037(1998).  
 CC -!- FUNCTION: PRODUCES ATP FROM ADP IN THE PRESENCE OF A PROTON  
 CC GRADIENT ACROSS THE MEMBRANE.  
 CC -!- SUBUNITS: F-TYPE ATPASES HAVE 2 COMPONENTS, CF(1) - THE CATALYTIC  
 CORE - AND CF(0) - THE MEMBRANE PROTON CHANNEL. CF(1) HAS FIVE  
 CC SUBUNITS: ALPHA(3), BETA(3), GAMMA(1), DELTA(1), EPSILON(1). CF(0)  
 CC HAS THREE MAIN SUBUNITS: A, B AND C.  
 CC -!- SUBCELLULAR LOCATION: CHLOROPLAST THYLAKOID MEMBRANE.  
 CC -!- SIMILARITY: BELONGS TO THE ATPASE EPSILON CHAIN FAMILY.  
 CC

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DR EMBL: X12889; CAA31381.1; -.  
 DR PF00401; ATP synt; DE; 1.  
 DR ATP synthesis; Chloroplast; Thylakoid membrane; CF(1);  
 KW Hydro-lase; Hydron ion transport.  
 KW SEQUENCE: 132 AA; 14472 MW; D826F274 CRC32; SQ

Query Match Score 43; DB 1; Length 132;  
 Best Local Similarity 66.7%; Pred. No. 2.53e+01;  
 Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;  
 SQ

Db 43 AVDIGILTI 51  
 QY 1 AAGIGILTV 9

CC

RESULT 10  
 ID SGAA\_HYPME STANDARD PRT; 404 AA.  
 AC 008374  
 DT 15-DEC-1998 (Rel. 37, Created)  
 DT 15-DEC-1998 (Rel. 37, Last sequence update)  
 DE SERINE--GLYOXYLATE AMINOTRANSFERASE (EC 2.6.1.45) (SGAT).  
 GN SGAA.  
 OS Hypomicrobium methylavorum.  
 OC Bacteria; Proteobacteria; alpha subdivision; Hyphomicrobium.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=GM2;  
 RX MEDLINE: 97054583.  
 RA HAGISHITA T., YOSHIDA T., IZUMI Y., MITSUNAGA T.;  
 RT "Cloning and expression of the gene for serine-glyoxylate  
 aminotransferase from an obligate methylotroph Hyphomicrobium  
 methylavorum GM2.";  
 RT Eur. J. Biochem. 241:1-5(1996).  
 CC -!- CATALYTIC ACTIVITY: L-SERINE + GLYOXYLATE = 3-HYDROXYPYRUVATE +  
 CC GLYCINE.  
 CC -!- COFACTOR: PYRIDOXAL PHOSPHATE.  
 CC -!- PATHWAY: SERINE PATHWAY.  
 CC -!- SIMILARITY: BELONGS TO CLASS-V OF PYRIDOXAL-PHOSPHATE-DEPENDENT  
 CC AMINOTRANSFERASES.  
 CC

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 CC

CC

RESULT 11  
 ID YXCC\_BACSU STANDARD PRT; 461 AA.  
 AC PA6333; 032289;  
 DT 01-NOV-1995 (Rel. 32, Created)  
 DT 15-JUL-1999 (Rel. 38, Last sequence update)  
 DT 15-DEC-1999 (Rel. 39, Last annotation update)  
 DE HYPOTHETICAL METABOLITE TRANSPORT PROTEIN IN IOLOS-HTPG INTERGENIC REGION.  
 GN YXCC OR SS92BR.  
 OS Bacillus subtilis.  
 OC Bacterium; Firmicutes; Bacillales/Clostridium group;  
 RN Bacillus/Staphylococcus group; Bacillus.  
 RP SEQUENCE FROM N.A.  
 RC SPTRAIN=168 / BGSC1A;  
 RX MEDLINE; 96093926.  
 RA YOSHIDA K.-I., SEKI S., FUJIMURA M., MIWA Y., FUJITA Y.;  
 RT "Cloning and sequencing of a 36-kb region of the Bacillus subtilis  
 genome between the gnt and iol operons.";  
 RL DNA Res. 2:61-69(1995).  
 RN [2]  
 RP REVISIONS.  
 RA SHIBAYAMA T., ISHTO I., AOYAMA D., YOSHIDA K.-I.;  
 RL Submitted (JUN-1997) to the EMBL/GenBank/DDBJ databases.  
 RN [3]  
 RP SEQUENCE FROM N.A.  
 RC SPTRAIN=168;  
 RA KUNST F., OGASAWARA N., YOSHIIWA H., DANOUCHI A.;  
 RL Submitted (NOV-1997) to the EMBL/GenBank/DDBJ databases.  
 CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (PROBABLE).  
 CC -!- SIMILARITY: BELONGS TO THE SUGAR TRANSPORTER FAMILY.  
 CC

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 CC

CC

RESULT 10  
 ID BAA21604\_1; -.  
 AC 008374  
 DT 15-DEC-1998 (Rel. 37, Created)  
 DT 15-DEC-1998 (Rel. 37, Last sequence update)  
 DE SUBTILIS; BG11360; YXCC.  
 DR PROSITE: PS00216; SUGAR TRANSPORT\_1; 2.  
 DR PROSITE; PS00217; SUGAR TRANSPORT\_2; 1.  
 DR HYPO; P00083; sugar\_irr; 1.  
 DR HYPOTHETICAL protein; Transport; Transmembrane.  
 FT TRANSMEM 15 35 POTENTIAL.  
 FT TRANSMEM 39 59 POTENTIAL.  
 FT TRANSMEM 77 97 POTENTIAL.



VCAD LAMBD STANDARD; PRT; 110 AA.  
 AC P03712;  
 DT 21-JUL-1986 (Rel. 01, Created)  
 DT 21-JUL-1986 (Rel. 01, Last sequence update)  
 DT 21-JUN-1994 (Rel. 29, Last annotation update)  
 DE HEAD DECORATION PROTEIN (GPD) (MAJOR CAPSID PROTEIN D).  
 GN D.  
 OS Bacteriophage lambda.  
 OC Viruses; dsDNA viruses, no RNA stage; Tailed phages; Siphoviridae;  
 Lambda phage group.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RX MEDLINE; 83189071.  
 RA SANGER F., COULSON A.R., HONG G.F., PETERSEN G.B.;  
 RT "Nucleotide sequence of bacteriophage lambda DNA.";  
 RL J. MOL. BIOL. 162:729-773(1982).  
 RN [2]  
 RP SEQUENCE.  
 RX MEDLINE; 84207913.  
 RA WITKIEWICZ H., SCHWEIGER M.;  
 RT "The head protein D of bacterial virus lambda is related to  
 eukaryotic chromosomal proteins.";  
 RT EMBO J. 1:1559-1564(1982).  
 CC -1- FUNCTION: STABILIZES THE HEAD SHELL FOLLOWING THE REARRANGEMENT  
 OF THE GPE SUBUNITS OF THE HEAD SHELL LATTICE THAT ACCOMPANIES  
 EXPANSION OF THE HEAD. THERE ARE APPROXIMATELY 420 COPIES OF  
 PROTEIN D PER MATURE PHAGE.  
 CC -1- SIMILARITY: TO BACTERIOPHAGE 21 HEAD DECORATION PROTEIN.  
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration  
 between the Swiss Institute of Bioinformatics and the EMBL outstation -  
 the European Bioinformatics Institute. There are no restrictions on its  
 use by non-profit institutions as long as its content is in no way  
 modified and this statement is not removed. Usage by and for commercial  
 entities requires a license agreement. (See <http://www.isb-sib.ch/announce/>  
 or send an email to license@isb-sib.ch).  
 CC DR EMBL; AAA90539; AA04349; J02459;  
 DR PIR; A04349; VHPDPL.  
 DR A23206; A23206.  
 KW Coat protein.  
 SQ SEQUENCE 110 AA; 11572 MW; FDD50011 CRC32;

Query Match 15 Score 75.0%; Best Local Similarity 55.6%; Pred. No. 4.15e+01;  
 Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Db 56 GAANGILAV 64  
 :|::|:||:|:  
 QY 1 AAGIGILTV 9

RESULT 15  
 ID FLA2\_METVO STANDARD; PRT; 216 AA.  
 AC P27804; P17602;  
 DT 01-AUG-1990 (Rel. 15, Created)  
 DT 01-AUG-1992 (Rel. 23, Last sequence update)  
 DT 15-DEC-1999 (Rel. 39, Last annotation update)  
 DE FLAGELLIN B2 PRECURSOR.  
 GN FLAB2.  
 OS Methanococcus voltae.  
 OC Archaea; Euryarchaeota; Methanococcales; Methanococcaceae;  
 Methanococcus.  
 RN [1]  
 RP SEQUENCE FROM N.A.  
 RC STRAIN=PS;  
 RX MEDLINE; 92041608.  
 RA KALMOKOFF M.L., JARRELL K.F.;  
 RT "Cloning and sequencing of a multigene family encoding the flagellins  
 of Methanococcus voltae."  
 RT J. Bacteriol. 173:7113-7125(1991).  
 RN [2]  
 RP SEQUENCE OF 13-32.

#cross-references MUID:93076110  
 #accession JQ1674  
 #molecule\_type DNA  
 ##residues 1-942 ##label CHA  
 ##cross-references GB:LO0670; NID:9166887; PID:AAA32876.1; PID:9166888  
 CLASSIFICATION #superfamily Protein kinase Xa21; leucine-rich alpha-2-glycoprotein repeat homology; protein kinase homology  
 KEYWORDS ATP; autophosphorylation; glycoprotein; phosphotransferase; receptor; serine/threonine-specific protein kinase; tandem repeat; transmembrane protein

FEATURE 1-22 #domain signal sequence #status predicted #label SIG  
 #product protein kinase TMK1 #status predicted #label MAT  
 23-942 #domain leucine-rich alpha-2-glycoprotein repeat  
 65-88 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR1\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR2\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR3\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR4\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR5\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR6\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR7\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR8\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR9\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR10\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR11\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR12\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR14\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR15\\  
 #domain leucine-rich alpha-2-glycoprotein repeat  
 homology #label LRR16\\  
 #domain transmembrane #status predicted #label TMN\\  
 594-602 #domain protein kinase homology #label KIN\\  
 86,99,158,164,171,363,533,587 #region protein kinase ATP-binding motif\\  
 586-872 #binding site carbohydrate (Asn) (covalent) #status predicted  
 616,634,717,719 #active-site Lys, Glu, Asp, Lys #status predicted  
 #length 942 #molecular\_weight 102387 #checksum 2851

SUMMARY 9.98; Score 88; DB 1; Length 942;  
 Best Local Similarity 37.5%; Pred. No. 8 34e-01; Gaps 1;  
 Matches 9; Conservative 10; Mismatches 4; Indels 1; Gaps 1;

Query Match 9.98; Score 88; DB 1; Length 942;  
 Best Local Similarity 37.5%; Pred. No. 8 34e-01; Gaps 1;  
 Matches 9; Conservative 10; Mismatches 4; Indels 1; Gaps 1;

Db 491 GLISFL-IGLVFCWKKRQKRF 513  
 Qy 31 GLITVILGVLLIGWCRRNGY 54

RESULT 15 YRHUR2 #type complete  
 ENTRY #molecule-type DNA #label YOK  
 TITLE dopachrome Delta-isomerase (EC 5.3.3.12) precursor - human  
 ALTERNATE\_NAMES dopachrome conversion factor; dopachrome oxidoreductase; dopachrome tautomerase; tyrosinase-related protein 2  
 ORGANISM #formal\_name Homo sapiens #common\_name man  
 DATE 13-Jan-1995 #sequence\_revision 17-May-1996 #text\_change  
 11-Jun-1999  
 ACCESSIONS S43510; S41089; I53786  
 REFERENCE S43510  
 #authors Yokoyama, K.; Suzuki, H.; Yasumoto, K.; Tomita, Y.;  
 Shishibata, S.  
 #journal Biochim. Biophys. Acta (1994) 1217:317-321  
 #title Molecular cloning and functional analysis of a cDNA coding  
 for human DOPACHROME tautomerase/tyrosinase-related  
 protein-2.  
 #cross-references MUID:94198295  
 #accession S43510  
 #molecule-type mRNA  
 #residues 1-519 #label YOK  
 #cross-references EMBL:DI7547; NID:9484259; PID:BA004484.1;  
 REFERENCE S41089  
 #authors Bouchard, B.; del Marmol, V.; Jackson, I.J.; Cherif, D.;  
 Dubertret, L.  
 #journal Eur. J. Biochem. (1994) 219:127-134  
 #title Molecular characterization of a human  
 tyrosinase-related protein-2 cDNA. Patterns of expression  
 in melanocytic cells.  
 #cross-references MUID:94139684  
 #accession S41089  
 #molecule-type mRNA  
 #residues 1-519 #label BOU  
 #cross-references GB:S69231; NID:9545618; PID:9545619  
 REFERENCE A44749  
 #authors Pawelek, J.M.  
 #journal Biochem. Biophys. Res. Commun. (1990) 166:1328-1333  
 #title Dopachrome conversion factor as an isomerase.  
 #contents annotation; reaction description  
 REFERENCE I53786

RESULT 14 F71010 #type complete  
 ENTRY hypothetical protein PH1380 - Pyrococcus horikoshii  
 ORGANISM #formal\_name Pyrococcus horikoshii  
 DATE 14-Aug-1998 #sequence\_revision 14-Aug-1998 #text\_change  
 14-Aug-1998  
 ACCESSIONS F71010  
 REFERENCE A71000  
 #authors Kavarabayasi, Y.; Sawada, M.; Horikawa, H.; Halkawa, Y.;  
 Hino, Y.; Yamamoto, S.; Sekine, M.; Baba, S.; Kosugi, H.;

```

#authors Cassady, J.L.; Sturm, R.A.
#journal Gene (1994) 143:295-298
#title Sequence of the human dopachrome tautomerase-encoding TRP-2
#DNA.

#cross-references MUID:94266170
#accession I53786
#status translated from GB/EMBL/DDBJ

##molecule_type mRNA
##residues 1-519 #!label RES
##cross-references GB:L18967; NID:939581; PIDN:AAA20870.1; PID:9399582
##experimental_source melanoma cell line A2058

GENETICS
#gene GDB:DCT; TYRP2
##cross-references GDB:231628; OMIM:191275
#map_position 13q32-13q32

FUNCTION
#description catalyzes the isomerization between 2-carboxy-1,2,3,
5-tetrahydroindolinium (dopachrome tautomer) and 5,
6-dihydroxyindole-2-carboxylic acid

PATHWAY
melanin biosynthesis
#superfamily monophenol monooxygenase
#copper; glycoprotein; intramolecular oxidoreductase;
#isomerase; melanin biosynthesis; transmembrane protein

CLASSIFICATION
copper; monophenol monooxygenase
FEATURE
#domain signal sequence #status predicted #label SIG\
1-23 #product dopachrome Delta-isomerase #status predicted
24-519 #label MAT
#domain transmembrane #status predicted #label TRM\
475-493
#binding-site carbohydrate (Asn) (covalent) #status
170,237,300,342, 377 predicted \
#binding-site copper (His) #status predicted \
#binding-site copper (His) #status predicted \
#binding-site copper (His) #status predicted \
#length 519 #molecular-weight 59145 #checksum 989

SUMMARY
Query Match 9.8%; Score 87; DB 1; Length 519;
Best Local Similarity 35.3%; Pred. No. 1.16e+00;
Matches 12; Conservative 11; Mismatches 10; Indels 1; Gaps 1;

Db 477 MGTVVALVGLFLVLLAFLQYRRRLRGYPALMETHL 510
Qy 30 IGILTVILGVLLIGCWTCRR-RNGYRALMDKSL 62

Search completed: Fri May 5 21:46:28 2000
Job time : 111 secs.

```

#journal Sugano, S. #title Genomics (1998) 49:458-461  
Cloning, expression analysis, and chromosomal localization of the cadherin  
superfamily.  
#cross-references MUID:98277460  
#accession T00041  
#status preliminary; translated from GB/EMBL/DBDJ  
#molecule\_type DNA  
##residues 1-1072 ##label YOS  
##cross-references EMBL:AB006756; NID:di184678; PID:di026123  
##experimental\_source clone BH-Pcdh-b

GENETICS  
#map\_position 4p15  
SUMMARY #length 1072 #molecular-weight 116462 #checksum 9727  
Query Match Score 99; DB 2; Length 1072;  
Best Local Similarity 50.0%; Pred. No. 1.88e-02;  
Matches 14; Conservative 8; Mismatches 4; Indels 2; Gaps 2;  
Organism Date 07-May-1999 #sequence\_revision 22-Jan-1999 #text\_change

Db 887 GIMTVLILIVVMARYCRSKNKGYEA 914  
Qy 31 GILTVLGVLLIGCWCR-R-RNGYRA 56

RESULT 6  
ENTRY T00042 #type complete  
TITLE BH-protocadherin PCDH7 (clone BH-Pcdh-c) - human  
ORGANISM #formal\_name Homo sapiens #common\_name man  
DATE 22-Jan-1999 #sequence\_revision 22-Jan-1999 #text\_change

ACCESSIONS T00042  
REFERENCE Z14074  
#authors Sugano, K.; Yoshitomo-Nakagawa, K.; Seki, N.; Sasaki, M.;  
#cross-references MUID:98277460  
#accession T00042  
#status preliminary; translated from GB/EMBL/DBDJ  
#molecule\_type mRNA  
##residues 1-1200 ##label YOS  
##cross-references EMBL:AB006757; NID:di184679; PID:di026124  
#experimental\_source clone BH-Pcdh-c

GENETICS  
#map\_position 4p15  
SUMMARY #length 1200 #molecular-weight 130337 #checksum 7152  
Query Match Score 99; DB 2; Length 1200;  
Best Local Similarity 50.0%; Pred. No. 1.88e-02;  
Matches 14; Conservative 8; Mismatches 4; Indels 2; Gaps 2;

Db 840 GIMTVLILIVVMARYCRSKNKGYEA 867  
Qy 31 GILTVLGVLLIGCWCR-R-RNGYRA 56

RESULT 7  
ENTRY C71600 #type complete  
TITLE rfrin PFB1020w - malaria parasite (Plasmodium falciparum)  
ORGANISM #formal\_name Plasmodium falciparum  
DATE 13-Nov-1998 #sequence\_revision 13-Nov-1998 #text\_change  
07-May-1999  
ACCESSIONS C71600  
REFERENCE Gardner, M.J.; Tettelin, H.; Carucci, D.J.; Cummings, L.M.;  
Aravind, L.; Koontz, E.V.; Shallow, T.; Yu, K.;  
Fujii, C.; Pederson, J.; Shen, K.; Jing, J.; Aston, C.;  
Lai, Z.; Schwartz, D.C.; Pertea, M.; Salzberg, S.; Zhou,  
L.; Sutton, G.C.; Clayton, R.; White, O.; Smith, H.O.;  
Fraser, C.M.; Adams, M.D.; Venter, J.C.; Hoffman, S.L.

#journal Science (1998) 282:1126-1132  
Chromosome 2 sequence of the human malaria parasite  
Plasmodium falciparum.  
#cross-references MUID:39021743  
#accession C71600  
#status preliminary; nucleic acid sequence not shown  
#molecule\_type DNA  
##residues 1-304 ##label GAR  
##cross-references GBAE001433; GB:AE001362; NID:g3845336; PID:g3845337;  
TIGR:PFB1020w  
##experimental\_source clone 3D7

GENETICS  
#gene PFB1020w  
SUMMARY #length 304 #molecular-weight 34000 #checksum 4789  
Query Match Score 96; DB 2; Length 304;  
Best Local Similarity 45.5%; Pred. No. 5.44e-02;  
Matches 15; Conservative 9; Mismatches 6; Indels 3; Gaps 3;  
Organism Date 22-Jan-1999 #sequence\_revision 22-Jan-1999 #text\_change

Db 263 EPCGIAAVLVLAVVYLJLYWYRRKNSYK 295  
Qy 26 EAAGIGILTV-TLGVLVLLIGC-W-YCRRRGYR 55

RESULT 8  
ENTRY T00043 #type complete  
TITLE BH-protocadherin-a - mouse  
ORGANISM #formal\_name Mus musculus #common\_name house mouse  
DATE 22-Jan-1999 #sequence\_revision 22-Jan-1999 #text\_change

ACCESSIONS T00043  
REFERENCE Z14075  
#authors Yoshihda, K.  
#submission submitted to the EMBL Data Library, August 1997  
#accession T00043  
#status preliminary; translated from GB/EMBL/DBDJ  
#molecule\_type mRNA  
##residues 1-1069 ##label YOS  
##cross-references EMBL:AB006758; NID:di127200; PID:di1033562

GENETICS  
#gene Pcdh7  
#map\_position 5C3-D  
SUMMARY #length 1069 #molecular-weight 116313 #checksum 4821  
Query Match Score 96; DB 2; Length 1069;  
Best Local Similarity 50.0%; Pred. No. 5.44e-02;  
Matches 14; Conservative 8; Mismatches 4; Indels 2; Gaps 2;

Db 887 GIMTVLILIVVMARYCRSKNKGYEA 914  
Qy 31 GILTVLGVLLIGCWCR-R-RNGYRA 56

RESULT 9  
ENTRY I37202 #type complete  
TITLE B-CAM protein - human  
ORGANISM #formal\_name Homo sapiens #common\_name man  
DATE 12-Aug-1996 #sequence\_revision 12-Aug-1996 #text\_change

ACCESSIONS I37202  
REFERENCE I37202  
#authors Campbell, T.G.; Foulkes, W.D.; Senger, G.; Trowsdale, J.;  
Garin-Chesa, P.; Rettig, W.J.;  
Cancer Res. (1994) 54:5761-5765  
#journal  
#title Molecular cloning of the B-CAM cell surface glycoprotein of  
epithelial cancers: a novel member of the immunoglobulin  
superfamily.  
#cross-references MUID:95042297  
#accession I37202  
#status preliminary; translated from GB/EMBL/DBDJ  
#molecule-type mRNA  
##residues 1-588 ##label RES



Title: >US-09-267-439-2  
Descriptor: (1-118) from US09267439.pep  
Target Score: 88  
Agreement: 1  
MPREDAHFTYGYPKKGCHS  
NADPDAYEKTSIAFOSPPDDVSD 118

scoring table: PAM 150  
Gap 11  
searched: 142080 seqs, 47172406 residues

post-processing: Minimum Match 0%

Listing first 45 summaries  
of 62

- 1:pir1 2:pir2 3:pir3 4:pir4

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

## Query result

No.	Score	Match	Length	DB	IB	Description	Pred.	No.
-----	-------	-------	--------	----	----	-------------	-------	-----

```

#journal Lurquin, C., Szikora, J.P.; Renauld, J.; Boon, T.
#title J. Exp. Med. (1994) 180: 35-42
#text A new gene coding for a differentiation antigen recognized by autologous cytolytic T lymphocytes on HLA-A2 melanomas [see comments].
#cross-references MUID:94275389
#accession I38506
#status preliminary; translated from GB/EMBL/DBJ
#molecule-type mRNA
##residues 1-118 ##label RES
##cross-references EMBL:006654; NID:9517022; PID:9517023
GENETICS
#gene GDB:MLANA
##cross-references GDB:358979
#map-Position 17q21-17q24
#length 118 #molecular-weight 13157 #checksum 3535
SUMMARY
Query Match 100.0%; Score 889; DB 2; Length 118;
Best Local Similarity 100.0%; Pred. No. 5.10e-179;
Matches 118; Conservative 0; Mismatches 0; Indels 0; Gaps 0
Db 1 MPREDAHFITYGPRKGHGSYTTAEEAAGIGLTIVLGVLLIGCWYRRRNGYRALMDK 60

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QY 1 MREDAHFIYGPKKKGHGSYTAAEAGIGLTVIQLIGWYRRRNGYRALMDK 60  
 molecular cloning, chromosome assignment and cell surface expression. MUID:87276135

Db 61 SLHVGTCALTRCPQEGFDHRSKVSQKEQNCEPYYVNPAPAYEKLSAEQSPPYSP 118  
 #cross-references S02293 accession not compared with conceptual translation

Qy \* 61 SLHVGTCALTRCPQEGFDHRSKVSQKEQNCEPYYVNPAPAYEKLSAEQSPPYSP 118  
 #molecule\_type mRNA  
 ##residues 1-127, 'M', '129-174, 'N', '176-191, 'M', '193-344 #label SEW  
 ##cross-references EMBL:Y00023; NID:950346; PIDN:CA68258.1; PID:950347

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 TITLE CD2 antigen precursor - mouse  
 ORGANISM #formal\_name Mus musculus #common\_name house mouse  
 DATE 02-Jul-1996 #sequence\_revision 02-Jul-1996 #text\_change  
 ACCESSIONS I49585  
 REFERENCE I49585  
 #authors Yagita, H.; Okumura, K.; Nakuchi, H.  
 #journal J. Immunol. (1988) 140:1121-1126  
 #title Molecular cloning of the murine homologue of CD2: Homology of the molecule to its human counterpart T11.  
 #cross-references MUID:88140313  
 #accession I49585  
 #status preliminary; translated from GB/EMBL/DDJB  
 #molecule\_type mRNA  
 ##residues 1-184 #label RES  
 #cross-references GB:MI834; NID:9192486; PIDN:AAA37397.1; PID:9309158  
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 SUMMARY #length 344 #molecular-weight 38337 #checksum 4699  
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 Best Local Similarity 40.4%; Pred. No. 1.88e-02;  
 Matches 21; Conservative 9; Mismatches 18; Indels 4; Gaps 4; Result 4  
 ACCESSIONS T00040 #type complete  
 REFERENCE 214074  
 #authors Yoshida, K.; Yoshitomo-Nakagawa, K.; Seki, N.; Sasaki, M.; Sugano, S.  
 #journal Genomics (1998) 49:458-461  
 #organism BH-protocadherin PCDH7 - human  
 #title Cloning, expression analysis, and chromosomal localization of BH-protocadherin (PCDH7), a novel member of the cadherin superfamily  
 #cross-references T00040  
 #accession T00040  
 #status preliminary; translated from GB/EMBL/DDJB  
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 #map\_position 4p15  
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 Matches 14; Conservative 8; Mismatches 4; Indels 2; Gaps 2;  
 ACCESSIONS T00041 #type complete  
 REFERENCE 214074  
 #authors Clayton, L.K.; Sayre, P.H.; Reinherz, E.L.  
 #journal Proc. Natl. Acad. Sci. U.S.A. (1988) 85:1615-1619  
 #title Exon-intron organization and sequence comparison of human and murine T11 (CD2) genes.  
 #cross-references MUID:88144486  
 #accession B28957  
 #molecule\_type mRNA  
 ##residues 1-344 #label DIA  
 #cross-references GB:MI19807; NID:9192479; PIDN:AAA37393.1; PID:9387122;  
 GB:J03622; GB:J03623  
 ##note the authors translated the codon TAT for residue 99 as Thr  
 #cross-references MUID:88004738  
 #accession S01347  
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 ##residues 1-127, 'M', '129-174, 'N', '176-190, 'NM', '193-344 #label CLA  
 #cross-references EMBL:X06143; NID:954223; PIDN:CAR29500.1; PID:g54224  
 REFERENCE S02293  
 #authors Sewell, W.A.; Brown, M.H.; Owen, M.J.; Fink, P.J.; Kozak, C.A.; Crumpton, M.J.  
 #journal Eur. J. Immunol. (1987) 17:1367-1370  
 #title Murine and human T11 (CD2) cDNA sequences suggest a common signal transduction mechanism.  
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